

ANTARCTIC DISPERSAL ROUTES, WANDERING CONTINENTS, AND THE ORIGIN OF AUSTRALIA'S NON-PASSERIFORM AVIFAUNA

By PAT VICKERS RICH *

* The Museum, Texas Tech University, Lubbock, Texas, U.S.A. 79409; present temporary address, The National Museum of Victoria, Russell Street, Melbourne, Victoria.

Introduction

In 1858, P. L. Sclater recognized Australia on the basis of its living avifauna, as a unique biogeographic unit, distinct from the Oriental fauna that characterized the Asian mainland. These two faunas complexly intermingle on the islands of the Indonesian archipelago, a situation reflected not only by the birds but by other vertebrate and invertebrate groups as well. A. R. Wallace in a number of papers (1863, 1869, 1876) initially described the avifaunas he encountered in this transitional zone; his work was followed by numerous refinements (Stresemann, 1927-34, 1939-41; Rensch, 1931) culminating with those which led Mayr (1941, 1944a-b, 1945a-b, 1972) to conclude that the major part of Australia's avifauna was derived from Southeast Asia. Although Darlington (1957) agreed with this thesis and Gentilli (1949) hinted at a dual origin, Serventy (1972, 1973) was the first to seriously consider multiple geographic origins of the Australian avifauna in light of the accumulating geological evidence suggestive of dynamic, not static, relationships of the continents during the Mesozoic and Cenozoic. Australia had not always been a neighbour of Asia. Serventy concluded, however, that excepting a few Australian families which might have used an Antarctic dispersal route, Mayr was correct in assuming that Southeast Asia had spawned most of the Australasian avifauna. Cracraft (1972, 1973) strongly supported an austral dispersal route for the ratites, megapodes, and possibly some parrots and suboscine passeriforms but otherwise subscribed to the thesis of northern origin for most other avian groups.

In the following paper, I wish to investigate the initial assumptions and the reasoning that have been used in previous studies to determine the source area for each of the Australian non-

passeriform families. In this way, I hope to clarify the certainties and uncertainties that accompany such determinations for each family, and to estimate which dispersal route (Antarctic or Indomalaysian) seems most probable for initial dispersal of each non-passeriform family between Australia and the remaining world. In order to evaluate the availability of the two routes for avian dispersal throughout the Mesozoic and Cenozoic, I have summarized: (1) data regarding the timing of break-up and separation of those continental plates closely associated with Australia during that time period; (2) paleoclimatological data available for the Antarctic and Indomalaysian dispersal routes, as well as for Australia during post-Paleozoic time; and (3) data available on phylogenetic relationships, world-wide diversity, endemism, distribution, and palaeontological record of all the non-passeriform avian families having a Cenozoic record in Australia.

Dispersal Routes To and From Australia

Geological Evidence

Antarctic Dispersal Route (Operative, Mesozoic and early Cenozoic)

Geophysical data collected primarily in the last decade strongly support the idea that a southern dispersal route between Australia and certain other Gondwana continents was present during much of the Mesozoic, and between Australia and some southern continents as late as the early Tertiary (Allan, 1969; Creer, 1970; Francheteau and Sclater, 1969; Smith and Hallam, 1970; and others). Such a route was most likely of an archipelagic nature (similar to the linkage between Asia and Australia today) as geologic data suggest that complete terrestrial continuity probably has not existed between South America, West

Antarctica, East Antarctica,¹ and New Zealand (Tedford, 1973a; Jardine and McKenzie, 1972) during the mid to late Mesozoic and Cenozoic.

Time of the break-up of the many continental segments, originally part of Gondwanaland, can be estimated by using several types

¹ Present-day Antarctica can no longer be considered a single, terrestrially continuous unit, but is instead composed of at least two segments, East and West Antarctica, each with a distinct geologic history (Hamilton, 1964, 1967; Dalziel and Elliot, 1971, 1973; Elliot, 1972). At present West Antarctica is comprised of a number of islands separated from the continental East Antarctica by basins with sub-sea level depths.

of geologic evidence. Despite the controversies that still attend the Africa-Madagascar, Madagascar-India, Africa-India, Australia-India, East-West Antarctica continental pairs, general agreement exists among most supporters of plate tectonic theory on the time of fragmentation of many of Gondwanaland's several segments. Such fragmentation did not occur simultaneously, however, and original rupture between continent-pairs may have begun long before actual terrestrial continuity between land masses was lost (e.g. East Africa at present). Thus, a distinction needs to be made between evidence indicating initial rifting or break-up (B in Table 1), and that indicative of active

TABLE 1

Evidence available for timing the break-up and dispersal of Gondwanaland during the Mesozoic and Cenozoic (see Figure 1 for summary). B, break-up occurring (involving displacements of less than 100 km.); S, large-scale separation of continental masses underway.

<i>Continental Pair</i>	<i>Time of Break-up/Dispersal</i>	<i>Evidence</i>
1. SOUTH AMERICA-AFRICA	S Post-Triassic to mid-Cretaceous (7, 13, 14, 17, 37, 38)	Polar wander curves.
	B Aptian (27, 39)	Salt deposits in coastal West Africa and coastal Brazil.
	B 106-136 m.y. (late Jurassic-early Cretaceous) (7, 8)	Occurrence of massive tholeiitic basalt flows dated by K/A method on both sides of Atlantic; Serra Geral Fm. of Brazil; Kaoko volcanics of Southwest Africa; Lupata volcanics of Africa.
	S 70-75 m.y. (26, 29, 46)	Magnetic banding (oldest anomaly, 31).
	B Post early Cretaceous (10)	Non-marine ostracod faunas virtually identical (at specific level) in Brazil and West Africa.
	S Albian (mid-Cretaceous) (10, 18, 28)	Fully marine conditions indicated by sediments over broad areas on either side of South Atlantic.
2. SOUTH AMERICA-WEST ANTARCTICA (probably archipelagic during Jurassic to recent)	S Late Turonian (12, 27, 28)	Fully marine conditions along western Africa. Ammonite faunas of North and South Atlantic similar.
	S > 20-25 m.y. but > late Mesozoic (22)	Magnetic banding in the Drake Passage for minimum date; similarity of tectonostratigraphic provinces until end of Andean Orogeny (late Cretaceous-early Tertiary) for maximum date.
	S Post-Cretaceous (24)	Paleomagnetism of volcanics and intrusives on Antarctic Peninsula indicate major bending of Peninsula occurred post-Cretaceous.
3. WEST ANTARCTICA-NEW ZEALAND (probably archipelagic, Jurassic to Recent)	S 80 m.y. (4, 5, 23, 26) (late Cretaceous)	Magnetic banding between Campbell Plateau and West Antarctica; oldest anomaly, 32.

<i>Continental Pair</i>	<i>Time of Break-up/Dispersal</i>	<i>Evidence</i>
	S 78 m.y. (19) (late Cretaceous)	Beginning of activity on Endeavour Fault (fault active between 47-78 m.y.).
	B Mid-Jurassic (3, 10)	Volcanics. Tasmanian dolerites, Ferrar dolerites (Antarctica).
4. EAST ANTARCTICA-AUSTRALIA (probably archipelagic Jurassic to Recent)	S Mid-Cretaceous-Tertiary (3, 13)	Polar wander curves.
	S 45-50 m.y. (1, 2, 4, 6, 10, 14, 16, 19, 26)	Magnetic banding between Australia and East Antarctica; oldest anomaly, 18.
	S Mid-late Eocene (1, 9, 10, 30)	Marine transgression in southern Australia; open marine limestones.
	S Late Oligocene (33, 36)	Erosional unconformity (marine) evidencing establishment of circum-Antarctic current.
	B Mid-Jurassic (42)	Dolerites in Tasmania and East Antarctica.
	B Late Jurassic to Cenomanian (43)	Rifting in southern Australia.
5. EAST ANTARCTICA-AFRICA	B Triassic-Jurassic (11)	Onset of rifting, southern Mozambique.
	B Mid-Triassic to mid-Jurassic 200-155 m.y. (8, 10, 20)	Volcanics. Extrusion of Stormberg, Karoo, Swaziland basalts. Intrusives in Rhodesia, South America.
	S Post mid-Jurassic-pre mid-Cretaceous (13)	Polar wander curves.
	S Pre late Cretaceous 140 m.y. (2, 12)	Magnetic banding between Africa and East Antarctica.
	S Pre Neocomian (15) (early Cretaceous)	Marine sediments.
	S Pre Valanginian (10) (early Cretaceous)	Marine sediments.
6. EAST ANTARCTICA-INDIA	S Mid-Jurassic-mid-Cretaceous (13, 14, 21)	Polar wander curves.
	S Pre-Cenomanian (10) (late Cretaceous)	Marine sediments between Antarctica and India.
	B Albian, 100-105 m.y. (early Cretaceous) (14, 21)	Volcanics. Extrusion of Rajmahal Traps.
7. AFRICA-MADAGASCAR (probably archipelagic Cretaceous to Recent)	S Mid-Jurassic-mid-Cretaceous (14)	Polar wander curves.
	B ?Permian (35)	Rifting.
	B Early Jurassic (32)	Volcanics.
	S Jurassic-late Cretaceous (31); pre early Cretaceous (45)	Volcanics, rifting, sediments (by Turoonian open marine conditions between Madagascar and Africa).
	B Cretaceous (11)	Rifting in East Africa.
	S Late Mesozoic (10)	Faunal.
	S Late Cretaceous (10)	Volcanics present in East Africa and Madagascar.
	S Late Eocene (34)	Marine sediments in Mozambique Channel.
8. INDIA-MADAGASCAR	S Post mid-Cretaceous (14)	Polar wander curves for India and Madagascar.
	S 70-75 m.y. (41)	Magnetic anomalies; oldest anomaly, 30.
	S Campanian (10) (late Cretaceous)	Marine sediments between India and Madagascar.
	B Late Cretaceous-early Tertiary, 59-65 m.y. (10, 14, 21, 40)	Volcanics. Deccan Traps of western India.
	S Post late Cretaceous (10)	Marine faunas (see over).

<i>Continental Pair</i>		<i>Time of Break-up/Dispersal</i>	<i>Evidence</i>
9. INDIA/MADAGASCAR- AFRICA	S	Mid-Jurassic- mid-Cretaceous (10)	Polar wander curves.
	S	70-75 m.y. (41)	Magnetic anomalies; oldest anomaly, 30.
	S	Pre latest Cretaceous, 67 m.y. (15)	Magnetic banding in NW sector of Indian Ocean.
	S	40 m.y. (26)	Magnetic banding.
	S	Post late Cretaceous (10)	Faunas (marine invertebrates) of Coro- mandel Coast of India quite similar to those of Assam and South Africa in late Cretaceous but distinct from those from Narbada Valley in NW India. Faunas of NW India most similar to those of North Africa and Europe. Evidence suggests non-marine barrier between northern and southern Africa, eastern and western India.
10. AUSTRALIA-INDIA	B	Late Jurassic-early Cretaceous, 100-105 m.y. (9, 12, 14, 15)	Volcanics. Rajmahal Traps (NE India), Banbury (Perth Basin) and Ashmore Reef (Aust.) basalts, Broom Beds (tuffs, Canning Basin, Aust.).
	S	Early Cretaceous (9, 12, 15)	Marine sediments. First occurrence of marine sediments (during Mesozoic) in Perth Basin. Thick sequence of non- marine sediments as young as Jurassic in subsurface, southern Perth Basin.
	S	Pre early Tertiary (3)	Sediments on either side of Ninetyeast Ridge in Indian Ocean 'show a unified history and little or no relative motion since early Tertiary'.
	S	Pre mid to upper Oligocene (33, 36)	Submarine erosion occurring in Tasman sea evidenced by thinness or absence of Oligocene sediments in this area and the scoured oceanic surface below such sediments; mid to upper Oligocene sedi- ments, when, present often intensely current-bedded, suggesting presence of submarine current.
	B	Latest Carboniferous- earliest Cretaceous (43)	Rifting in Western Australia.
	B	Permian-Neocomian (42)	Vulcanism, Banbury Basalt, latest Neo- comian tholeiite.
	S	Late Jurassic (in NW part of Australia) to late Neocomian-Aptian (in SW Australia) (43)	Magnetic banding in Indian Ocean and initiation of marine transgression in Western Australia.
	S	Pre latest Jurassic (Tithonian) (44)	Sediments (oldest) overlying oceanic crust near Western Australian coast.
11. AUSTRALIA- NEW ZEALAND (probably archipelagic Cretaceous to Recent)	S	80 m.y. (4)	Magnetic banding between Campbell Plateau and W. Antarctica, Australia, and E. Antarctica indicates New Zealand moving north while Australia still attached to E. Antarctica.
	B	Paleocene-Eocene	Volcanics. Beginning of significant vol- canic activity in eastern Australia that continue through much of the Tertiary.
	?	51 m.y. (3)	

The following papers are referred to in Table 1 under their respective numbers:

1. Jones, 1971
2. LePichon and Heirtzler, 1968
3. Wellman, McElhinny, and McDougall, 1969
4. Griffiths and Varne, 1972
5. Pitman, Herron and Heirtzler, 1968
6. Weissel and Hayes, 1971
7. Creer, 1970
8. Dietz and Holden, 1970
9. Veevers, 1971
10. Smith and Hallam, 1970
11. Sowerbutts, 1972
12. Tarling, 1971
13. Francheteau and Sclater, 1969
14. McElhinny, 1970
15. Veevers, Jones, and Talent, 1971
16. McElhinny and Wellman, 1969
17. Vilas and Valencio, 1970
18. Burke, Dessauvage, and Whiteman, 1971
19. Christoffel and Ross, 1970
20. Dietz and Sproll, 1970
21. McDougall and McElhinny, 1970
22. Dalziel and Elliot, 1971
23. Griffiths, 1971
24. Hamilton, 1964
25. Maxwell, *et al.*, 1970
26. Heirtzler, *et al.*, 1968
27. Reyment, 1969
28. Funnell and Smith, 1968
29. Dickson, Pitman, and Heirtzler, 1968
30. McGowran, 1971 (ms)
31. von der Borch, 1972
32. Flores, 1970
33. Kennett, *et al.*, 1972
34. Simpson, *et al.*, 1972
35. Kent, 1972
36. Carter and Landis, 1972
37. Creer, Embleton, and Valencio, 1969
38. Larson and LaFountain, 1970
39. Leyden, Bryan, and Ewing, 1972
40. Wellman and McElhinny, 1970
41. McKenzie and Sclater, 1971
42. Harrington, *et al.*, 1973
43. Veevers and Evans, 1973
44. Heirtzler, *et al.*, 1973
45. Green, 1972
46. Valencio and Vilas, 1972
47. Jones, 1972
48. Dingle, 1973

occurrence of magnetic anomalies in the intervening ocean basins, (3) the first differences in the polar wander paths of segments previously part of one continental plate, and (4) differences in the tectonostratigraphic provinces of two previously geologically similar and contiguous continental plates (see Smith and Hallam, 1970; and particularly Tarling and Tarling, 1971 for more detail on the dating methods mentioned above). Such data evidencing break-up and dispersal of the Gondwana continents since the Triassic are summarized in Table 1 and Figure 1 for each continental pair, except East Antarctica-West Antarctica (see discussion below). Figure 1 specifically indicates the variety of the data used to estimate the time of break-up of each continental pair, while Table 1 details that data, indicating whether break-up or active dispersal was underway.

Fragmentation within Gondwanaland began as early as the Triassic, but actual severance in terrestrial continuity between those parts of the southern continents connected in the late Palaeozoic (approx. 225 m.y. B.P.) was a late Mesozoic and even early Tertiary phenomenon (approx. 70-130 m.y. B.P.; see figures 1-2, 7-10, and Table 1). The first complete separation was apparently between Africa and East Antarctica in the late Jurassic or early Cretaceous (Heirtzler, *et al.*, 1968, Francheteau and Sclater, 1969; Dietz and Sproll, 1970; Smith and Hallam, 1970; Dietz and Holden, 1970; Dingle and Klinger, 1971; Heirtzler and Burroughs, 1971; Tarling, 1971; Sowerbutts, 1972; Jones, 1972; Dingle, 1973). Although not so well understood, the severance of any close alliance between India and Australia probably occurred in the early Cretaceous (Heirtzler, *et al.*, 1968; Veevers, 1971; Tarling, 1971; Veevers, Jones, and Talent, 1971; Crawford, 1971²), certainly not later than late Mesozoic (von der Borch, *et al.*, 1972). Thus the shortest dispersal route between Australia and the remaining continents during the late Mesozoic and much of the Paleogene was across Antarc-

² Crawford argues that during the Phanerozoic India was closely apposed to Africa with a seaway separating India and Australia. Just how broad such a water gap would have been is uncertain at present.

separation of continental masses (S in Table 1) with concomitant loss of non-marine dispersal pathways. Widespread occurrence of vulcanism and extensional faulting, sometimes accompanied by evaporite deposits, can be the first indication of continental break-up. Similarly, active separation between continental blocks is evidenced by: (1) the first widespread occurrence of marine sediments after a long period of continental sedimentation, (2) the first

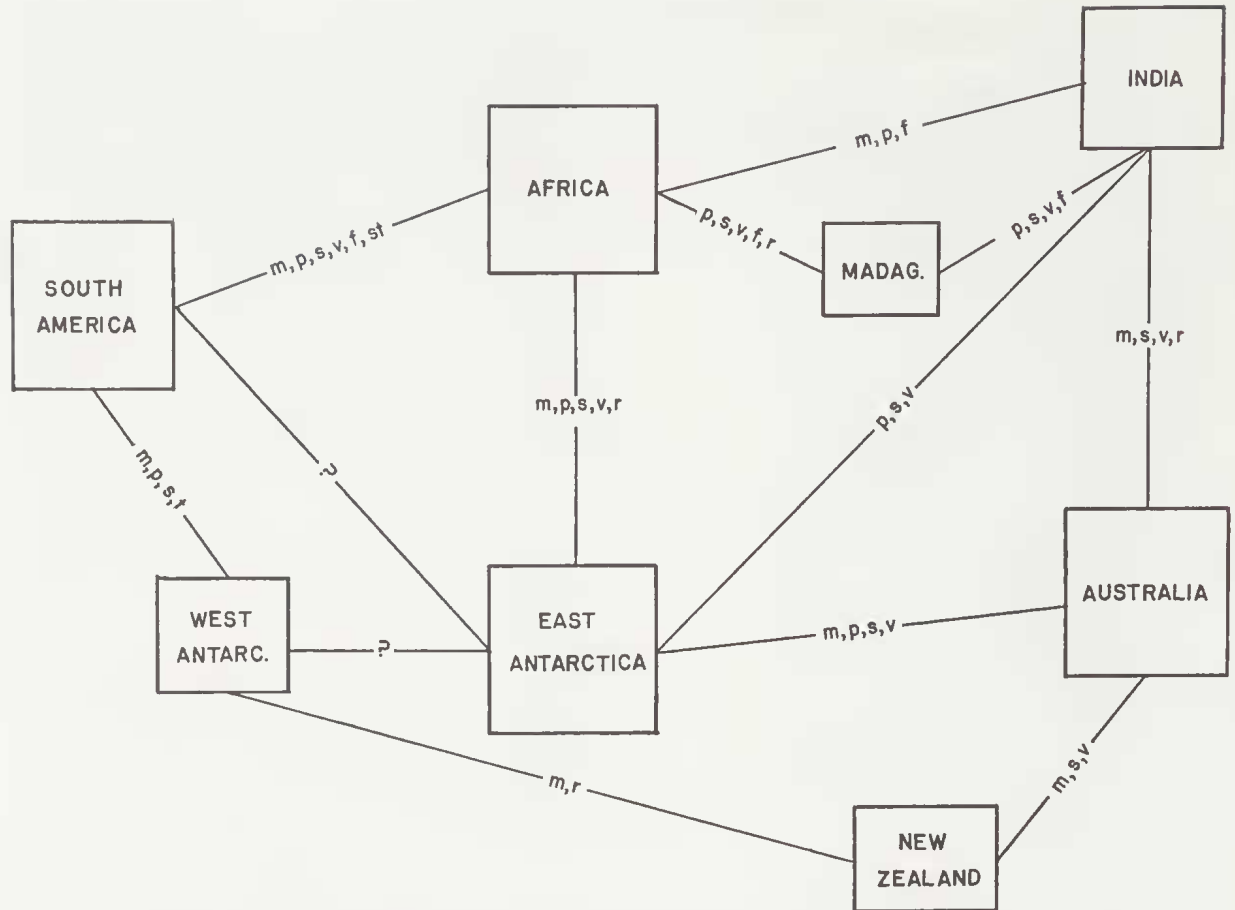


Fig. 1—Evidence available for timing the post-Paleozoic break-up and dispersal of Gondwanaland. The following symbols were used: m, magnetic anomalies on the seafloor; p, polar wander curves; s, widespread occurrence of marine sediments after long period of non-marine sedimentation and other sedimentological data; v, occurrence of widespread volcanism; f, faunal (both marine and terrestrial) similarities and differences; r, onset of major extensional rifting; t, similarity of tectonostratigraphic provinces; st, occurrence of widespread salt deposits.

tica. Asia, at this time, was as much as 30-50° north of Australia (Dietz and Holden, 1970; Tedford, 1973a; Jardine and McKenzie, 1972; Raven and Axelrod, 1972).

In the late Mesozoic (see figure 4), East Antarctica and Australia seemingly formed a continuous land mass (Jones, 1971; Le Pichon and Heirtzler, 1968; Smith and Hallam, 1970; McElhinny, 1970; Heirtzler, et al., 1968; Veevers, 1971; McGowran, 1971). Connection

at this time to South America, as to New Zealand, appears to have been archipelagic. Similarly a West Antarctic archipelago may have lain between South America and New Zealand (Fleming, 1962; Tedford, 1973a; Elliot, 1972). Yet unresolved and critical to late Mesozoic and Cenozoic reconstructions, however, is the precise relationship of West Antarctica to South America and East Antarctica. Many workers suggest that during much of the Mesozoic West Antarctica and New Zealand were all part of one plate's compressive margin (Elliot, in press, 1972), and must have developed in close proximity to a continental sediment source, East Antarctica and Australia (van der Linden, 1969), with the southern Andean Cordillera and the Antarctic Peninsula, although not necessarily a linear chain (Dalziel and Elliot, 1971, 1973), forming a link between these two continents. Severe disruption of this perhaps already archipelagic

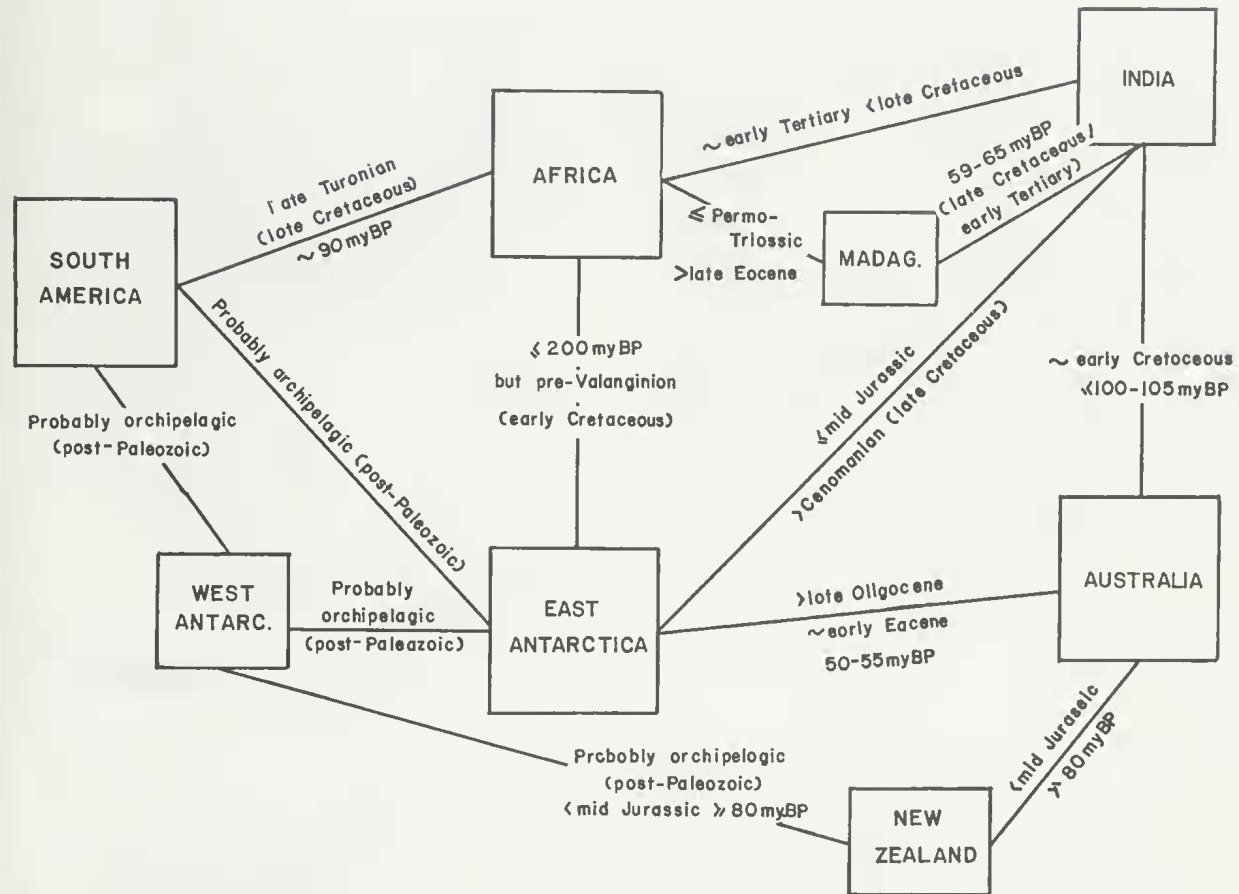


Fig. 2—Estimate of time of most recent severance of land connections between presently distinct continental masses, once part of Gondwanaland.

cordillera may have begun prior to 82 m.y. B.P. (Dalziel and Elliot, 1971, 1973) with transform or transcurrent faulting rotating the Ellsworth Mountains (now situated between East Antarctica and the Antarctic Peninsula) (Dalziel and Elliot, 1971, 1973; Elliot, 1972; Schopf, 1969). During the late Mesozoic and early Cenozoic, South America and West Antarctica may have approached one another more closely than at present (Dalziel and Elliot, 1973). Some workers, however, suggest that East and West Antarctica were independent entities during the Mesozoic, becoming closely apposed only in the Cenozoic (Beck, 1972). Others further suggest that the post-Jurassic tectonic histories of southern South America and West Antarctica have been decidedly independent

and that deformation of the Scotia Arc occurred prior to the late Mesozoic (Katz, 1973). Elliot (pers. comm., 1974) has pointed out that this does not necessarily mean that insular linkage was lacking between the two areas during the Mesozoic and part of the Cenozoic, however. More general agreement attends the positioning of New Zealand decidedly closer to Marie Byrd Land in West Antarctica until about 80 million years ago, when it apparently began its northward drift with the foundering of the Campbell Plateau and opening of the Tasman Sea (Griffiths, 1971; Griffiths and Varne, 1972). The opinion of most geologists dealing with these areas, however, despite the differing details of their reconstructions, is that any dispersal route involving West Antarctica from mid-Mesozoic to the Recent would probably have been archipelagic (Elliot, 1972; Jardine and McKenzie, 1972; Dalziel, pers. comm., 1973).

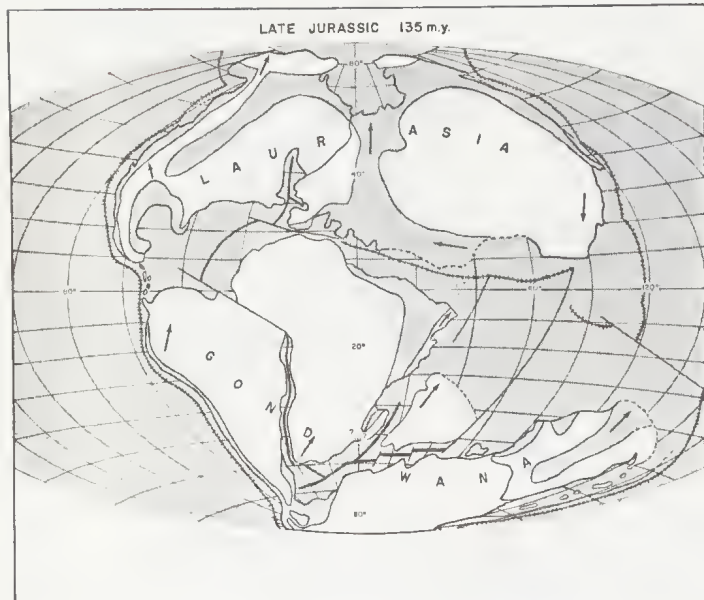
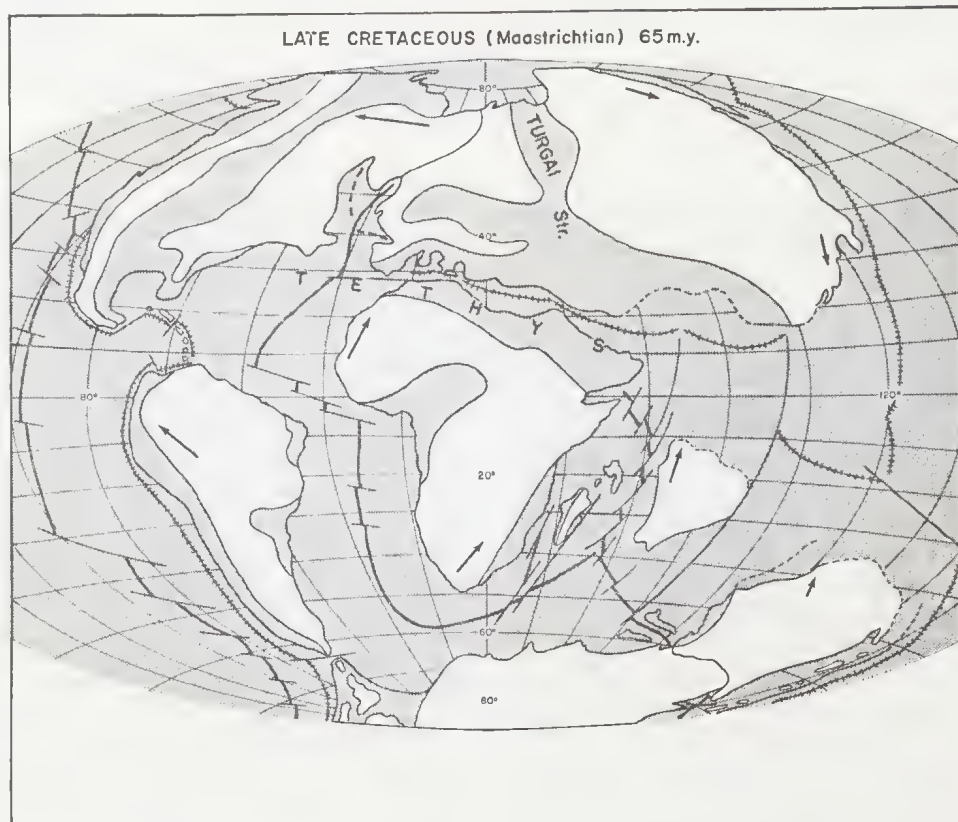


Fig. 3—Continental arrangement during the late Jurassic (135 m.y.) as modified from Tedford, 1973a. In such reconstructions, the earth's surface is divided into a series of crustal plates that move relative to each other. Three types of boundaries separate these plates: (1) transform faults (light lines) where the plates slide laterally past one another; (2) ridges (dark lines), where new molten rock is added to the edges of two plates and they subsequently move away from one another; and (3) trenches (hatched lines), where one plate slides beneath another. Arrows indicate direction of relative motion.

Fig. 4—Continental arrangement during the late Cretaceous (65 m.y.) as modified from Tedford, 1973a.



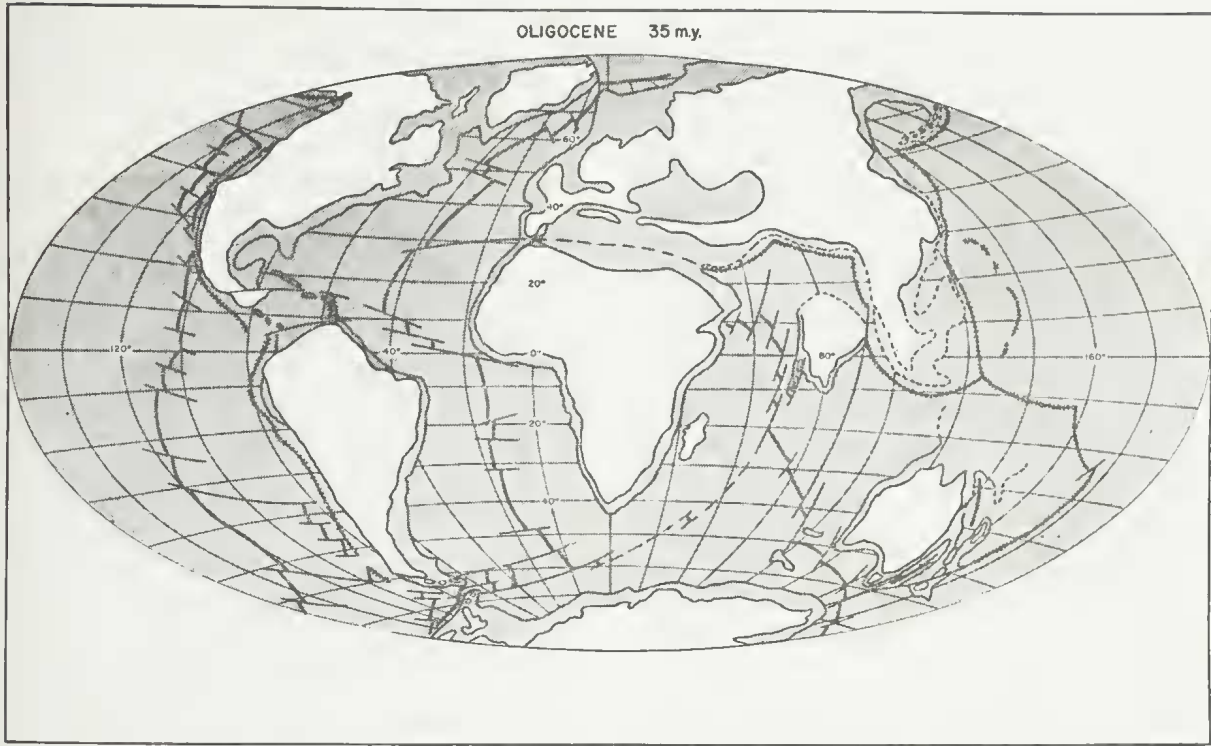
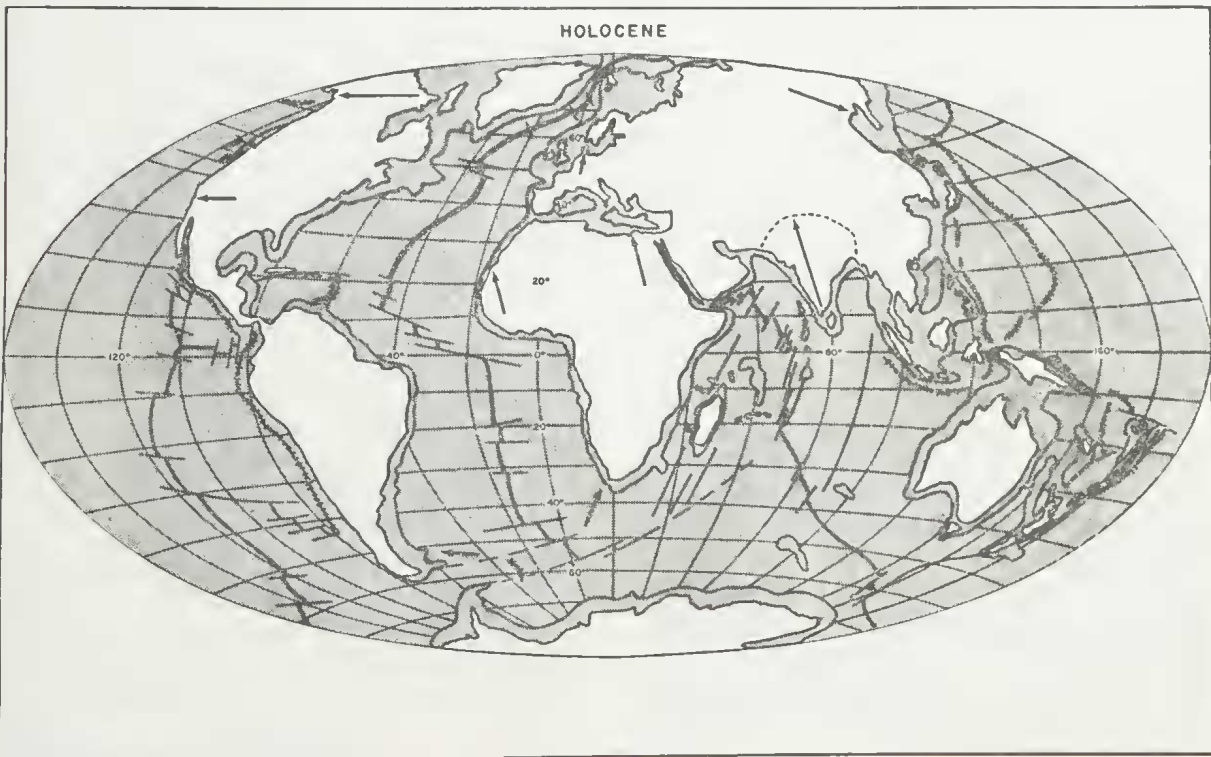


Fig. 5—Continental arrangement during the Oligocene (35 m.y.) as modified from Tedford, 1973a.

Fig. 6—Continental arrangement at present.



Only in the Tertiary, the early to mid-Eocene (50-55 m.y. B.P.), and certainly by the late Oligocene (Kennett, *et al.*, 1972), did Australia sever its connection with East Antarctica (Le Pichon and Heirtzler, 1968; Jones, 1971; Veevers, 1971; Smith and Hallam, 1970; Davies and Smith, 1971; McGowran, 1971; Vogt and Conolly, 1971; Griffiths and Varne, 1972; Weissel and Hayes, 1972) and begin its northward journey, which is still in progress (see figure 9). From a position of 60-70° south latitude (measured from Canberra) in the Cretaceous, Australia moved to within 10° of its present position (see figure 10) by the Miocene (Wellman, McElhinny, and McDougall, 1969; Veevers, Jones, and Talent, 1971; Raven and Axelrod, 1972). New Zealand continued separating from both Australia and West Antarctica, increasing the isolation that presently characterizes it. Thus, during the late Mesozoic and early Tertiary, the separation of Australia and New Zealand from their present Asian and most Indonesian neighbours was great with close apposition occurring only after the Oligocene (see Hamilton, 1972a-b; Haile and McElhinny, 1972; Stauffer and Gobbett, 1972; *contra* Ridd, 1971, and Audley-Charles *et al.*, 1972).

Indonesian Dispersal Route (Operative, Late Cenozoic)

The mid-Cenozoic close approach of Australia to Asia (including Southeast Asia, see Haile and McElhinny, 1972 and closely apposed Indonesia, see Ben-Avraham and Uyeda, 1973; Hamilton, 1972a-b), presented new possibilities for dispersal to and from Palaeo-antarctica. With Antarctica's movement south of the Australo-Antarctic Ridge and concomittant deterioration in its climate accompanying positioning closer to the pole, it was further isolated from Australia. Thus, its importance as a dispersal route declined as that of Indonesia increased. By the Miocene, the principal dispersal route for terrestrial organisms to and from Australia was the Indomalaysian Archipelago (McKenna, 1973), including the Philippines, much of which was subaerial by this time (Hamilton, 1972a; van Bemmelen, 1949; Grabau, 1926).

Throughout the Cenozoic the area between Southeast Asia and Australia has probably been dotted with islands, many of which have been ephemeral, occasionally disappearing down oceanic trenches or submerging along with older parts of ocean basins. In some senses a dispersal route of sorts probably existed across this archipelago in the Paleogene, but distances between islands as well as the total length of the route were much greater than at present and prohibitive in all but the rarest cases of non-marine vertebrate dispersal. Most of central and southern New Guinea, the leading edge of the Australian plate in the Paleogene, was underwater until the late Oligocene-early Miocene (Raven and Axelrod, 1972; Davies and Smith, 1971; Tedford, 1973a; Audley-Charles *et al.*, 1972). During the Miocene, a southward migrating island arc system collided with the northern edge of the Australian crustal plate (see figure 7 for explanation), adding the final, northern segment to form modern New Guinea (Hamilton, 1972b) and causing considerable uplift. By late Cenozoic, therefore, extensive emergent lowland areas and developing highlands were present in New Guinea. In the Indo-malaysian area (extremely complex geologically) much of the Malay Peninsula, and at least parts of Sumatra, Java, Borneo, and Sulawesi were emergent during the early Cenozoic (Grabau, 1926; van Bemmelen, 1949; Kummel, 1961). Major parts of the Andaman-Sumatra-Java-Timor-Outer Banda-Seram chain of islands, however, are composed of uplifted subduction melanges (a jumble of sediments, volcanics, and metamorphics typifying an area where one major plate of the earth's crust is underthrusting another) produced during the Miocene and uplifted sometime later (Hamilton, 1972a-b; Katili, 1971). During the late Pliocene, folding further uplifted many of the previously homogeneous lowland areas in both Indonesia and New Guinea (Kummel, 1961), providing a mountainous, temperate dispersal route (Raven and Axelrod, 1972) between Asia and Australia, probably for the first time in the Mesozoic-Cenozoic. Further emergence during the Pleistocene concurrent with development of ice sheets elsewhere in the world, shortened water gaps, and made the Indo-

malayan area a very probable route for dispersal of terrestrial organisms.

Paleoclimatological Evidence

Antarctic Dispersal Route

Based on paleobotanical data, the climatic regime of the Antarctic Peninsula and East Antarctica was much less severe during the late Mesozoic and early Tertiary than at present. During the Jurassic, scouring rushes (Equisetales), ferns (Filicales), cycadophytes (Cycadophyta), and conifers are known from several localities in Grahamland, West Antarctica (Plumstead, 1962, 1964; Bibby, 1966). Unfortunately, only one post-Triassic, pre-Tertiary flora is known from East Antarctica (Victoria Land); it contains a cycadophyte as well as three types of conifers very similar to those known from India, East Africa, and Australia in strata of early Jurassic age. Such assemblages in both East and West Antarctica suggest equable, moist conditions, warmer than at present, particularly in order to accommodate the cycadophytes, which were experiencing latitudinal restriction in the Northern Hemisphere during the Jurassic (Plumstead, 1964). The presence of freshwater actinopterygian fish in lower Jurassic sediments of East Antarctica (Schaeffer, 1972) lends corroborative evidence for more equable, certainly non-glacial Mesozoic conditions in parts of Antarctica. Mega and microfossil evidence indicates that at least two families of ferns (Cyatheaceae and Selaginaceae), several conifers (*Araucaria*, *Agathis*, *Podocarpus*) (Andrews, 1961) and angiosperms including the Southern Beech (*Nothofagus*), mustards (Cruciferae), Myrtaceae, Proteaceae, Loranthaceae, and possibly *Drimys* (Winteraceae) were growing on Seymour Island in West Antarctica during the Cretaceous (Cranwell, 1959). Likewise, floras very tentatively dated as early to mid-Tertiary from King George and Seymour Islands, also in West Antarctica, contain scouring rushes, ferns, conifers (including *Araucaria*), as well as a variety of angiosperms, including *Nothofagus* (Adie, 1953; Cranwell, 1959; Plumstead, 1964). Approximately ninety per cent of the

dicotyledons in these floras have non-entire margins and small leaves, more typically reminiscent of temperate plant species living in mild to cool conditions at present (Axelrod and Bailey, 1968). Again, the flora of East Antarctica is not as well known, with only two lower Tertiary localities: one in the McMurdo Sound area (Cranwell et al., 1960; McIntyre and Wilson, 1966) and a second near Prydz Bay (Kemp, 1972a-b) on the Australian side of East Antarctica. Much of the pollen from these localities is similar to that from West Antarctica. The McMurdo sample derived from Eocene erratic boulders and recycled sediments in the Ross Sea includes conifer, palm, *Podocarpus*, and *Nothofagus* pollen, among others. Probable *in situ* late Oligocene sediments in the Ross Sea have recently produced many of the same elements with *Nothofagus* and Podocarpaceae dominant and minor percentages of Proteaceae and Myrtaceae. Noticeably absent is pollen of groups presently found on subantarctic islands such as Gramineae, Cyperaceae, Umbelliferae, or Compositae (E. Kemp., pers. comm., 1974). The Prydz Bay pollen flora derived from Cretaceous-Eocene sediments is decidedly more varied, containing at least 21 genera including *Nothofagus* (*fusca*, *menziesii*, and *brassi*), *Podocarpidites*, *Gleicheniidites*, *Osmundacites*, and *Cyanthidites* (Filicales) reminiscent of dominant taxa in the early Tertiary Australian flora. Certainly as diverse a flora as this seems incompatible with a continent-wide glaciation prior to the Oligocene.

In summary, fossil floras from East and West Antarctica, although admittedly rare, suggest a temperate climate for parts of this region capable of supporting sizeable trees and a diverse assemblage of plants during much of the Mesozoic, possibly as late as the Miocene and even the Quaternary on the Falkland Islands, which support no forests today (Axelrod, 1960). Much of this same flora that occurred in Antarctica was also present in southern South America, Kerguelen Island, New Zealand, and parts of Australia (Axelrod, 1960) during the early to mid-Tertiary. This flora was apparently rather similar to the temperate and cool temperate rainforests of present-day southern Chile, Patagonia, south-

eastern Australia, Tasmania, New Zealand, and New Caledonia.

Additional information of paleoclimatic significance has been derived from geologic and fossil invertebrate data. Recent electron micrograph studies of sand grain surface features based on submarine core samples from the Subantarctic Pacific, has been interpreted to indicate that glaciation, although probably of limited extent, was underway in parts of Antarctica as early as the Eocene (Geitzenauer *et al.*, 1968; Gram, 1969; Margolis and Kennett, 1970, 1971a-b; Kennett and Brunner, 1973). Apparently glacially transported sand grains were rafted away from the Antarctic as part of icebergs and deposited as the ice melted. Accompanying the first occurrence of such sands, Mandra (1969, 1971; also in Mandra and Mandra, 1970) noticed a drop in silicoflagellate species diversity, which was interpreted as indicative of cooling of the southern oceans at that time. In the early and middle Oligocene cores, ice rafted sand is no longer a dominant feature, if present at all, and silicoflagellate diversity in such sediments is higher than in immediately older sediments, indicating warmer seas than in the earlier Tertiary, perhaps owing to the establishment of the circum-Antarctic current at about this time.

Additional geologic evidence mustered to suggest an Eocene glaciation in West Antarctica has been recovered from Marie Byrd Land. LeMasurier (1970) has noted the occurrence of hyaloclastites, volcanic rocks known to form at present only in subglacial or subaqueous eruptions of lava. Accumulations of hyaloclastites up to 400 metres thick with no evidence of subaerial lava flows have been interpreted to indicate ice at least that thick in some places during the Eocene, providing, of course, that such volcanics belong to a single flow and providing that Le Masurier is correct in assuming that this area in West Antarctica was above sea level when such lavas were extruded. Other hyaloclastite accumulations in the same area indicate glaciation from the Oligocene to the Pleistocene where ice thickness may have reached 1200 metres by the Pliocene. Lava flows dated as late Miocene in the Jones Mountains of West Antarctica are known to overlie

"consolidated glacial deposits and extensive areas of glacially scoured bedrock" (Newell, 1972, p. 64). Even though much of West Antarctica (particularly the Antarctic Peninsula) supported *Nothofagus* dominated forests perhaps as late as the Miocene, glaciation may have been underway in limited areas as early as the Eocene. The evidence for widespread early Tertiary glaciation, however, based on such sedimentological and volcanic data has recently been challenged by Mercer (1973), who believes the botanical evidence for both segments of Antarctica suggest nothing more than local glaciation until 3-4-6 m.y. B.P. Certainly by 4-5 m.y. B.P. glaciation on a continental scale is suggested by widespread erosion surfaces over much of the Ross Sea continental shelf, evidently produced by a grounded ice shelf (Hayes, *et al.*, 1973), but glacial history of Antarctica prior to this time has yet to be resolved.

Unfortunately, owing to the lack of post-Jurassic palcomagnetic data from East Antarctica (primarily because of the lack of post-Jurassic rocks cropping out above the present ice cover, as well as the paucity of samples even from West Antarctica) there is as yet no coherent picture of the South Pole position with respect to Antarctica during the late Mesozoic and early Tertiary. A number of papers (Francheteau and Sclater, 1969; McElhinny and Wellman, 1969; Crawford, 1971; McElhinny, 1970; among others) summarizing available paleomagnetic evidence derived from continents exclusive of Antarctica, including Australia, present no consensus on East Antarctica's mid to late Mesozoic or Early Tertiary latitudinal position. Data reported by McElhinny and Wellman (1969) and Francheteau and Sclater (1969) do not exclude the possibility of a high latitude (yet *outside* the Antarctic Circle) dispersal route along the Victoria Land and Wilkes Land coast of East Antarctica during the early Tertiary. Almost all reconstructions, however, place West Antarctica within the Antarctic Circle during late Mesozoic and Cenozoic, and yet the evidence of *Nothofagus*-dominated forests in this area is incontrovertible. Either such forests were able to withstand a yearly cycle with three months of

perpetual darkness or parts of East and West Antarctica lay at a lower latitude. This only points out the need for additional geophysical data before the latitudinal position of both major segments of Antarctica during the Mesozoic and Cenozoic is clear.

Indomalaysian Dispersal Route

In marked contrast to the cool, temperate floras of Paleogene Antarctica, are floras from the late Mesozoic and Cenozoic Indomalaysian area. Although no thorough, current summary of the Southeast Asian and Indonesian fossil floras is available, a number of early papers (Barbour, 1929; Colani, 1917, 1920; Edwards, 1923; Göppert, 1854; Holden, 1916; Kräusel, 1922, 1925, 1926; Seward, 1941; Andrews, 1961 among others) discussing leaf, wood, and fruit evidence and more recently papers (Chaney, *et al.*, 1963; Endo, 1964; Endo and Fujiyama, 1966; Germeraad, *et al.*, 1968; Jones *et al.*, 1966; Konno, 1966; Manten, 1969) describing the pollen, allow an estimation of the late Mesozoic and Cenozoic vegetation of this area. By the Cretaceous, several elements now characteristic of the warm, humid tropical lowlands of Southeast Asia and Indonesia were present there. Germeraad, *et al.* (1968) reported *Nypa* palm, now restricted to the mangrove swamps of the Indomalaysian region, from late Cretaceous sediments of Sarawak. Paleogene floras of northern Viet Nam (Colani, 1920), Borneo (Kräusel, 1925), Sumatra (Kräusel, 1922), and Java (Göppert, 1854) contain many of the tropical elements found in floras of these areas today: Palmae, Pandanaceae, *Ficus*, *Quercus* (closely related to tropical oaks in this genus), Piperaceae, Anacardiaceae or Annonaceae, Moraceae, Musaceae, Dipterocarpaceae, *Rhizophora* (mangrove), among others. Some of these early floras contain only tropical forms, but others exhibit a mixture of these and more temperate elements such as *Taxus*, *Phoebe*, *Betula*, *Ben-zoin* (Colani, 1920), characteristic of the Himalayas or more northern areas in Asia, and probably represent elements of upland floras of Indonesia and Southeast Asia. Miocene and younger floras from the Indomalay region are more diverse, containing a large

number of presently tropical taxa, with first occurrences of such forms as Sapindaceae, Sterculiaceae, and definite Anacardiaceae among others. To the north of Southeast Asia, however, cool to warm temperate assemblages had differentiated by the early Cenozoic. Floras of northern Thailand (Endo and Fujiyama, 1966), Siberia, Manchuria, Korea, and Japan (Barbour, 1929; Chaney, *et al.*, 1963; Endo, 1964; Endo and Fujiyama, 1966) during early to mid-Cenozoic, commonly contained such forms as *Sequoia*, *Taxodium*, *Pinus*, *Alnus*, *Populus*, Betulaceae, Ulmaceae, Platanaceae, Juglandaceae, Ericaceae as well as some families (e.g. Fagaceae, Aceraceae) held in common with the more southerly areas.

Undoubtedly, a detailed re-evaluation of much of the early work on Indomalaysian paleofloras must be carried out before the floral history of this area will be understood. However, with recent palynological investigation confirming many of the earlier identifications of Cenozoic tropical forms based on leaves, wood, and fruit, apparently several families of plants presently restricted to the tropics occurred in Southeast Asia as early as the Cretaceous and Paleogene. During the entire history of the modern avian families, and certainly during the Neogene, then, much of the Indomalaysian area probably supported a warm, humid tropical flora, not so different (at the family level, at least) from that occupying the same area today.

Australia

Instrumental in the development of the Australian avifauna were the dramatic changes in the climatic regime affecting Australia during post-Palaeozoic times. Raven and Axelrod (1972), Gill (1961), Dorman (1966), and Burbidge (1960) summarize much of the geochemical, floral, faunal, pedologic, and lithologic data that are employed to reconstruct Australia's past climate.

Most important as temperature indicators are radioisotope, paleobotanical, and faunal data. Paleotemperature determinations, based on oxygen isotope studies (O^{18}/O^{16} ratios) of Cretaceous belemnites from Australia and New Zealand (Stevens and Clayton, 1971), indicate

decreasing near shore water temperatures from an early Cretaceous (ca. 110 m.y. B.P.) high to an Albian-Cenomanian (ca. 100 m.y. B.P.) low (N.Z. samples: from 42° S. present latitude, ca. 16° C., comparable to averages N.Z. temperatures at 39-40° S.; Australian samples from Lake Eyre region at 28° S. present latitude, ca. 15° C., comparable to temperatures at 41° S. at present). Temperatures rose again in the Coniacian-Santonian (ca. 18° C. from samples taken at 42° S. present latitude in N.Z., comparable to temperatures at 36-37° S. at present) but began dropping again during the Campanian (ca. 70-75 m.y. B.P.), continuing into the Paleocene. Oxygen isotope studies on shells of several different molluscan genera (collected from localities in southern Victoria between the present 38-39° S. latitudes) suggest a continued drop in temperatures into the late Eocene (to ca. 12-16° C.) followed by a rise to a Cenozoic maximum in the mid-Miocene (22-25° C.), probably accompanying the onset of Australia's northward drift from Antarctica. Since the Miocene, temperatures have continued to drop, with minor fluctuations, to the present (12° C. for southern Victoria) (Dorman, 1966; Dorman and Gill, 1959a-b).

Cretaceous and early Tertiary floras indicate widespread temperate conditions evidenced by mixed forests of austral gymnosperms and evergreen angiosperms (Podocarpaceae, Araucariaceae, Araliaceae, Myrsinaceae, Protaceae, Winteraceae, Atherospermataceae, Epacridaceae, Malvaceae, Loranthaceae, Sapindaceae, Casuarinaceae, Myrtaceae, and *Nothofagus* (Fagaceae); Raven and Axelrod, 1972). During this time *Araucaria*, now restricted in Australasia to areas in New Guinea and coastal eastern Australia, where average temperature of the coldest month is 11° C., grew in Tasmania and southern Victoria. *Nothofagus brassi*, presently restricted to altitudes above 3000 feet in New Guinea, was widespread in the early Tertiary of southern and eastern Australia. Mid-Tertiary floras, on the other hand, indicate restriction of such temperate forests (*Nothofagus-Araucaria* dominated) to the southern and northern margins of the continent and the dominance of the *Eucalyptus-Acacia* assemblage

that contained such additional taxa as *Casuarina*, *Melaleuca*, *Hakea*, and *Eremophila*. Most of this mid to late Cenozoic flora appears to have been (and still is) adapted to the drier, more seasonal conditions that became prevalent on a continental scale as Australia moved into the high pressure "horse latitudes" with their accompanying xeric climates. Sclerophyll forest, savannas, and more arid vegetation covered much of Australia, while rain forests were present along the eastern, mountainous edge of the continent (Tedford, 1973a).

In New Guinea, with the emergence of extensive lowlands in the late Oligocene-early Miocene, a rich tropical flora with strong Malaysian affinities and quite distinct from that of mainland Australia gained a foothold. Only much later in the Pliocene and Pleistocene, accompanying the tectonism that produced extensive temperate, montane environments as well as xeric conditions in areas of rain shadow (Raven and Axelrod, 1972), was there much exchange between New Guinea and Australia. At this time, too, northern temperate forms (e.g. *Castanopsis*, *Rhododendron*) were able to enter New Guinea and Australia from Asia for the first time along the temperate uplands created by the mid to late Cenozoic tectonism.

Invertebrate faunas (particularly foraminifera, mollusca, and echinoids) indicate, by faunal composition and diversity, a Tertiary temperature maximum in the early Miocene.³ Only at this time were any of the larger foraminifera (*Trybliolipidina*, *Cycloclypeus*) present in the southern Australian Bass Strait Province. Likewise, the first major reef building episode in Australia since the Palaeozoic began in the Miocene when Old World reef communities, as well as other tropical invertebrates

³ The mid-Tertiary temperature rise in Australia, indicated by invertebrate faunas and oxygen isotope ratios, correlates well with the drift of this continent northward into lower, more tropical latitudes. Throughout most of the world, temperatures steadily (with minor fluctuations) declined throughout the Cenozoic. Once Australia neared its present position sometime in the Miocene and its northward drift slowed, its climate deteriorated in synchrony with the rest of the world, climaxing in the low temperatures of the Pleistocene.

(Palmiere, 1971), colonized the Australian continental shelf (Newell, 1971, 1972). In Victoria, many groups of marine molluscs (cones, cowries and volutes in particular) exhibited increased diversity in the early to mid-Miocene, as well as a marked increase in Indopacific taxa (Darragh, 1973; Darragh, 1974). By the late Miocene (Cheltenhamian), however, temperatures began dropping as indicated by the striking reduction both in number of individuals and in diversity of both molluscs and echinoderms. By Kalimnan times (about 6 m.y. B.P.), the marine invertebrate faunas in the continental shelf areas around Australia were essentially modern (Brown, Campbell, and Crook, 1968), indicating temperatures not so different from those today.

Vertebrate faunas, too, indicate changes in humidity and temperature in much of Australia during the Cenozoic. Lungfish, confined to a small area of southeastern Queensland in the late 19th century (Grigg, 1964), are present in Cretaceous sediments in southern Victoria (Schaeffer, 1969) and at a number of localities of Tertiary age in the Australian interior. Teleost fish are known from most of the localities producing lungfish, yet many of these areas no longer support permanent streams. Crocodylians, now rarely found further south than Rockhampton on the northeast coast and the Ord River (Gill, 1961) in the northwest, ranged into northern South Australia and Victoria in the Tertiary, demonstrating greater humidity and yearly minimum temperatures than encountered today in those areas. Diverse fossil avifaunas from the mid to late Tertiary of interior South Australia and the Northern Territory (see figures 3-4, table II, and later section of paper) include flamingos (Phoenicopteridae) and even aningas (Miller, 1963b, 1966a), the latter restricted to warm regions with permanent streams and abundant streamside vegetation today and probably for some time in the past (Rich, 1972, 1974). Fossil mammals, similarly, suggest a more humid environment during the Tertiary (see Tedford, 1973a for discussion of the fossil marsupial faunas of Australia). In interior South Australia and the Northern Territory a diverse assemblage of medium to large-sized mammals are known

from probable mid-Miocene sediments and include koalas, several taxa of diprotodontids (extinct marsupial family closely related to the kangaroos), wombats, phalangers, thylacoleonids, monotremes, and even porpoises among others (see Stirton, Tedford, and Woodburne, 1968; Tedford, 1973a-b). A number of smaller forms including dasyurids, phalangerids, and peramelids were also present. The occurrence of koalas, as well as the diversity of mammals in these mid-Tertiary faunas, suggests a more heavily vegetated, more humid environment during the mid-Tertiary than presently exists in these areas today, a suggestion certainly corroborated by other vertebrate as well as floral evidence.

Summary

During the Mesozoic and as late as mid-Eocene (ca. 50 m.y. B.P.), geological and paleobotanical data suggest that a southern dispersal route across East Antarctica and archipelagic West Antarctica was the most likely connection between Australia and other continents. Southeast Asia during this period was probably far to the north, perhaps as much as 30-50° of latitude. At least many of the islands making up West Antarctica, as well as parts of East Antarctica, supported substantial forests dominated by several kinds of gymnosperms, including *Araucaria* and *Podocarpus* as well as a number of angiosperms including *Nothofagus*. Part of this route, possibly along the Victoria Land and Wilkes Land coast of East Antarctica and appropriate islands in West Antarctica, may have lain north of the Antarctic circle, thus providing a route having some daylight hours throughout the entire year. By the end of the Miocene, Australia had severed its connections with East Antarctica and drifted north to within 10° of its present position, strengthening the dominance of the Indomalaysian dispersal route, which has been most influential during the latter part of the Cenozoic. Simultaneous with northward drift, climatic changes affecting the Australian continent brought about a shift from the early Cenozoic cool temperate flora dominated by proteads and *Nothofagus* to the mid to late Cenozoic (including the present) zoned vegetation com-

posed of arid (in interior Australia), tropical (in New Guinea and Northern Australia), and cool temperate (in southeastern Australia and Tasmania) elements. Such marked changes in Australia's geographic position, its climate and floras, and its links with the rest of the world, undoubtedly have had a tremendous effect on its Cenozoic vertebrate faunas, including its birds.

Origin of the Non-passeriform Avifauna of Australia

Australian Avifaunal Composition

The living terrestrial and fresh-water non-passeriform avifauna of Australia and New Guinea is composed of some 44 families, of which the only endemics are the emu and cassowary (Casuariidae: *Dromaius* and *Casuaris*) and the collared hemipode (Pedionomidae) (see Table 3). If those birds restricted to the Australia-New Guinea-Southwest Pacific (excluding New Zealand) area are considered, then the megapodes (Megapodiidae) and owl frogmouths (Aegothelidae) can be added to the list of endemic families. In addition, a few distinct subfamilies are endemic to Australia and/or New Guinea and the Southwest Pacific, including the Cercopsinae, Loriinae, Cacatuinae, and Micropsittinae, all except the first, members of the Psittacidae (parrots). At the generic level, 34 taxa (exclusive of those in the above-mentioned families and subfamilies) are endemic to Australia and/or New Guinea alone (see Table 2) and an additional 28 are endemic to Australia and/or New Guinea and the Southwest Pacific (again excluding New Zealand). The fossil record adds two families not present in Australia today: the Dromornithidae, composed of large to truly gigantic ratite birds endemic to Australia, and the Phoenicopteridae (flamingos), both of which survived into the Pleistocene in Australia; flamingos, of course, yet survive in several parts of the world.

History of the Australian Non-Marine Avifauna (Figures 7-8)

Early Cretaceous (perhaps late Jurassic; J. Warren, pers. comm., 1974) lacustrine sedi-

ments in southeastern Victoria have produced feathers (Talent, *et al.*, 1966; Waldman, 1970) and record an early presence of birds on the Australian continent. These, together with a few Eocene-Oligocene penguin bones from southeastern Australia (Simpson, 1946, 1957, 1959, 1965, 1970; Gill, 1959a-b; Finlayson, 1938; Anonymous, 1959), a single record of an unidentified bird bone (now lost) from early Tertiary sediments near Sunnybank, southeastern Queensland (Hill and Denmead, 1960, p. 349; Houston, 1967, p. 85), and a feather impression as well as a possible forelimb of a bird from near Wannon, western Victoria (Chapman, 1910; David, 1950), constitute the early record of birds in Australia. By mid-Miocene, however, a moderately diverse assemblage of modern avian families was present, known from several localities (see Table 3) in the interior of Australia and includes:⁴

Dromornithidae (mihirung birds) (Rich, 1973)

Dromaiinae (emus) (Rich, 1973)

Pelecanidae (pelicans) (Miller, 1966b)

Phalacrocoracidae (cormorants) (Rich, 1973)

Accipitridae (an eagle) (Rich, 1973)

Anatidae (ducks) (Rich, 1973)

Phoenicopteridae (flamingos) (Miller, 1966a)

cf. Gruidae (cranes) (Stirton, *et al.*, 1968)

cf. Rallidae (rails) (Stirton, *et al.*, 1968)

Charadriiformes, undetermined family (shorebirds) (Rich, 1973)

Laridae (gulls) (Stirton, *et al.*, 1968)

Burhinidae (thick knees, stone curlews) (Stirton, *et al.*, 1968; Rich, 1973)

Caprimulgiformes (nightjars, frogmouths) (Rich, 1973).

The avifauna, even at this time, was strikingly modern with only the dromornithids (4-5 genera in the Miocene) and the flamingos (2 genera in the Miocene) forming an odd

⁴ To complete the list of Australian Miocene fossil birds, Wilkinson, H. E., 1969, reported an albatross (Diomedidae) from Victoria. Since this is a marine bird, however, it will not be considered in the above discussion.

TABLE 2

Endemics to Australia-New Guinea-Southwest Pacific (after Peters 1931-1945)

- † Australia and/or New Guinea
- ‡ Australia and/or New Guinea and Southwest Pacific
- * Australia and/or New Guinea, Southwest Pacific, Asia

FAMILY LEVEL

- † Casuariidae (including †*Casuarus* and †*Dromaius*)
- † Pedionomidae (†*Pedionomus*)
- ‡ Megapodidae (‡*Megapodius*, †*Leipoa*, †*Alectura*, ‡*Telegalla*, ‡*Aepyodius*)
- * Podargidae (**Podargus*)
- ‡ Aegothelidae (‡*Aegothales*)
- * Hemiprocnidae (**Hemiproctus*)

SUBFAMILY LEVEL

- † Ceropsinae (†*Cereopsis*)
- ‡ Loriinae (‡*Trichoglossus*, ‡*Pseuteles*, †*Glossopsitta*, ‡*Oropsitta*, †*Lathamus*, ‡*Chalcopsitta*, †*Pseudeos*, ‡*Domicella*, ‡*Charmosyna*, †*Oreopsittacus*, †*Neopsittacus*, ‡*Psittaculirostris*)
- ‡ Cacatuinae (‡*Probosciger*, †*Calyptorhynchus*, †*Callocephalon*, ‡*Cacatua*, †*Eolophus*, †*Nymphicus*)
- ‡ Microsittinae (‡*Micrositta*)

GENERIC LEVEL

- ‡ *Zonerodius* (Ardeidae)
- * *Xenorhynchus* (Ciconiidae)
- † *Anseranas* (Anatidae, Plectropterinae)
- † *Stictonetta* (Anatidae, Anatinae)
- † *Malacorhynchus* (Anatidae, Anatinae)
- ‡ *Salvadorina* (Anatidae, Nyrocinidae)
- ‡ *Chenonetta* (Anatidae, Anserinae)
- † *Biziura* (Anatidae, Oxyurinae)
- ‡ *Henicopernis* (Accipitridae, Elaninae)
- † *Lophoictinia* (Accipitridae, Milvinae)
- † *Haemirostra* (Accipitridae, Milvinae)
- * *Haliastur* (Accipitridae, Milvinae)
- † *Erithroriorchis* (Accipitridae, Accipitrinae)
- † *Harpyopsis* (Accipitridae, Buteoninae)
- ‡ *Synoicus* (Phasianidae)
- † *Anurophaps* (Phasianidae)
- ‡ *Eulabeornis* (Rallidae)
- * *Rallina* (Rallidae)
- ‡ *Tribonyx* (Rallidae)
- * *Amaurornis* (Rallidae)
- * *Poliolimnas* (Rallidae)
- † *Rallcula* (Rallidae)
- ‡ *Gymnocrex* (Rallidae)
- † *Megacrex* (Rallidae)
- † *Peltohyas* (Charadriidae)
- † *Cladorhynchus* (Recurvirostridae)
- * *Esacus* (Burhinidae)
- ‡ *Stiltia* (Glareolidae)
- ‡ *Procelsterna* (Laridae)
- ‡ *Gygis* (Laridae)
- * *Ptilinopus* (Columbidae, Treroninae)
- ‡ *Megaloprepia* (Columbidae, Treroninae)
- * *Ducula* (Columbidae, Treroninae)
- † *Lopholaimus* (Columbidae, Treroninae)
- † *Petrophassa* (Columbidae, Columbinae)

- * *Macropygia* (Columbidae, Columbinae)
- * *Chalcophaps* (Columbidae, Columbinae)
- † *Phaps* (Columbidae, Columbinae)
- † *Ocyphaps* (Columbidae, Columbinae)
- † *Lophophaps* (Columbidae, Columbinae)
- † *Geophaps* (Columbidae, Columbinae)
- † *Histriophaps* (Columbidae, Columbinae)
- † *Leucosarcia* (Columbidae, Columbinae)
- † *Geopelia* (Columbidae, Columbinae)
- ‡ *Gymnophaps* (Columbidae, Columbinae)
- ‡ *Reinwardtoena* (Columbidae, Columbinae)
- ‡ *Gallicolumba* (Columbidae, Columbinae)
- ‡ *Henicophaps* (Columbidae, Columbinae)
- † *Trugon* (Columbidae, Columbinae)
- ‡ *Otidiphaps* (Columbidae, Columbinae)
- ‡ *Caloenas* (Columbidae, Columbinae)
- ‡ *Goüra* (Columbidae, Goürinae)
- ‡ *Eclectus* (Psittacidae, Psittacinae)
- ‡ *Geoffroyus* (Psittacidae, Psittacinae)
- † *Polytelis* (Psittacidae, Psittacinae)
- ‡ *Aprosmictus* (Psittacidae, Psittacinae)
- ‡ *Alisterus* (Psittacidae, Psittacinae)
- ‡ *Platycercus* (Psittacidae, Psittacinae)
- † *Purpureicephalus* (Psittacidae, Psittacinae)
- † *Psephotus* (Psittacidae, Psittacinae)
- † *Neophema* (Psittacidae, Psittacinae)
- † *Melopsittacus* (Psittacidae, Psittacinae)
- † *Pezoporus* (Psittacidae, Psittacinae)
- † *Geopsittacus* (Psittacidae, Psittacinae)
- † *Psittichas* (Psittacidae, Psittacinae)
- † *Psittacella* (Psittacidae, Psittacinae)
- * *Loriculus* (Psittacidae, Psittacinae)
- ‡ *Scytlirops* (Cuculidae, Cuculinae)
- * *Ninox* (Strigidae)
- ‡ *Uroglaux* (Strigidae)
- * *Eurostopodus* (Caprimulgidae)
- ‡ *Dacelo* (Alcedinidae, Daceloninae)
- ‡ *Tanyssipectera* (Alcedinidae, Daceloninae)
- ‡ *Melidora* (Alcedinidae, Daceloninae)
- † *Clytoceyx* (Alcedinidae, Daceloninae)

element leaving no living representatives in Australia. Both of the latter groups survived into the Pleistocene but not the Recent.

By the Pliocene cassowaries (*Casuarus*) had differentiated and were present in New Guinea (Plane, 1967) while a third genus of flamingo, *Phoeniconaias*, occurred for the first and only time in Australia (*Phoeniconotius* is not known from sediments younger than Miocene, while *Phoenicopterus* persisted into the Pleistocene).

Pleistocene faunas add the first known Australian occurrences of the Podicepsidae (*Podiceps*, grebes), the Threskiornithidae (spoonbills), cf. *Cygnus atratus* (Black Swan), the Megapodiidae (mound builders, including a gigantic form nearly twice the size of any living megapode, from several localities in eastern Australia), the Phasianidae (quail), the Falconidae (falcons), the Turnicidae



Fig. 7—Tertiary and Mesozoic localities producing fossil birds in Papua-New Guinea and Australia: Awe (1); Bullock Creek (2); Riversleigh (3); Alcoota (4); Peak Downs (5); Chinchilla (6); Lake Kanunka (7); Lake Palankarina (8, 9); Eurinilla Creek and Lake Pinpa (10); Port Noarlunga (11); Christie's Beach (12); Pritchard Brothers' Quarry (13); Mt. Gambier (14); Devil's Den (15); Spring Creek (Minhamite) (16); Beaumaris (17); Endurance Pit (18); Lake Ngapakaldi (19); Lake Pitikanta (20); Redbank Plains (21); Kangaroo Well (22); Bugaldi (near Coonabarabran (23); Redruth, Wannan (24); and Koonwarra (25).

(*Turnix*, bustard quail), the Pedionomidae (plains wanderers), the Scolopacidae (sandpipers), the Chionididae (*Chionis*, sheathbill), the Columbidae (pigeons, doves), the Psittacidae (parrots and allies), the Tytonidae

(*Tyto*, barn owl), and several passeriforms. A number of additional groups reported by DeVis (1888a-b, 1889a-b, 1891a-c, 1905; in Gregory, 1906) and Longman (1945) need restudy to ascertain their taxonomic positions (Phalacrocoracidae, **Ciconiidae**, Anatidae, Rallidae, **Otididae**⁵).

Thus, at least 23 of the 44 families constituting the modern non-passeriform avifauna of Australia have mid to late Cenozoic histories there, while two additional families have Miocene to Pleistocene records but no modern representatives in the Australian avifauna. Most certainly the lack of a Tertiary record for many other families is owing to the small

⁵ Taxa named in boldfaced type indicate earliest known occurrence in Australia.



Fig. 8—Quaternary localities producing fossil birds in Papua-New Guinea and Australia: Diamantina River (no specific locality known) (1); Cassidy Locality (2); Warburton River Localities (3); Cooper's Creek Localities (4); Lake Callabonna (5); Thorbindah (6); Cudde Springs (7); Ashford Caves (Bone Cave) (8); Bingara (9); Canadian Lead (Gulgong) (10); Walli and Wellington Caves (11); Wombeyan Caves (Guineacor) (12); Lake Menindee (13); Lake Tandou (14); Baldina Creek (15); Normanville (Salt Creek) (16); Kangaroo Island (17); Henschke's Cave and Victoria Cave (18); Penola (19); Mt. Gambier (20); King Island (21); Madura Cave (22); Scott River (23); Mammoth Cave (24); Darling Downs (Kings Creek, Warwick) (25); Drover's Cave (26); Gore (27); Buchan Caves (28); Weeke's Cave (29); Brother's Island, Pt. Lincoln (30); and Lancefield (31).

number of Cenozoic vertebrate producing localities presently known in Australia.

Owing to the limited nature of the fossil record in Australia, first occurrences of avian families and genera have no special significance regarding which dispersal route was being used, or in what direction movement occurred. By the Miocene, when the first extensive avifaunas are known in Australia, Southeast Asia and Australia were close enough to one another for avifaunal interchange to have been underway. Unfortunately we know almost nothing of the earlier Australian terrestrial avifaunas, which could have marked significance regarding South American affinities, enhanced by the existence of an Antarctic dispersal in the early Tertiary.

TABLE 3

Summary of the Avifaunas from Australian Cenozoic Localities¹
(if reference not cited, from Rich, 1973)

TERTIARY
EOCENE

- Queensland Interbedded sediments in unnamed basalt overlying Darra Fm. and underlying Sunnybank Fm. at Sunnybank, Aves, undetermined (now lost). Cribb, *et al.*, 1960.
- South Australia *Christie's Beach*. 'Transitional Marl' Member, base of Blanche Point Marls, *Palaeudyptes cf. antarcticus* (Spheniscidae, penguin). Simpson, 1946, 1957.
- South Australia *Port Noarlunga Jetty*. Banded Marl Member of Blanche Point Marls, *Palaeudyptes cf. antarcticus*. Simpson, 1946, 1957.

OLIGOCENE

- South Australia *Mount Gambier*. Gambier Limestone. Spheniscidae (penguin). Simpson, 1946, 1957.
- South Australia *Pritchard Brothers' Quarry*. Gambier Limestone, 7½ miles WNW of Mt. Gambier, *Palaeudyptinae* (penguin). Simpson, 1946, 1957.

MIOCENE

- South Australia *Snake Dam Locality*. Mooloorina Station, cf. Dromornithidae (mihirung bird, egg shell). M. Woodburne, M. Archer, pers comm., 1973.
- South Australia *Lake Pinpa*. Unnamed unit; *Phalacrocorax* sp. (Phalacrocoracidae, cormorant), Anatidae (duck), Burhinidae (thick-knee, stone curlew).
- South Australia *Eurinilla Creek*. Unnamed unit; Anatidae (duck), cf. Phoenicopteridae (flamingo).
- South Australia *Lake Palankarina*. Etadunna Formation, Ngapakaldi fauna (including localities LFM Loc. 9, U.C.M.P. Locs. V-5762, V-5763, V-5764, V-5765, V-5770, V-5771, Mammalon Hill); *Pelecanus tirarensis* (Pelecanidae, pelican), Phalacrocoracidae or Anhingidae (cormorant or darter), Phoenicopteridae, gen. et sp. undet. (flamingo), *Phoeniconotius eyrensis* (Phoenicopteridae), (undet.), *Oxyura* (Anatidae, duck), cf. Gruidae (crane), Rallidae (rail), Burhinidae (thick-knee), Charadriiformes (shorebirds), Laridae (gull), Stirton, Tedford, and Woodburne, 1967.
- South Australia *Lake Pitikanta*. Etadunna Formation, Ngapakaldi fauna (including Discovery Basin U.C.M.P. V-5774 and U.C.M.P. V-6150); *Phoenicopterus novaehollandiae* (Phoenicopteridae, flamingo). Miller, 1963b.
- Queensland *Riversleigh*. Carl Creek Limestone, Riversleigh fauna; Dromornithidae (new gen., new sp.) (mihirung bird).
- Victoria *Devil's Den* (Glenelg River). Glenelg Group, Balcomian Stage, north of Dartmoor; *Anthropodyptes gilli* (Spheniscidae, penguin). Simpson, 1959.
- South Australia *Kangaroo Well*. Unnamed formation, Kangaroo Well fauna, Aves, undetermined. Stirton, Tedford, and Woodburne, 1967.
- South Australia *Lake Ngapakaldi, Leaf Locality*. Wipijiri Formation, Kutjamarpu fauna (U.C.M.P. V-6213); *Dromaius* (Casuariidae, emus), Dromornithidae (mihirung bird); *Ngapakaldi 2* (U.C.M.P. V-6213), Dromornithidae. Stirton, Tedford, and Woodburne, 1967.
- Northern Territory *Bullock Creek*. Camfield Beds, Bullock Creek fauna; Dromornithidae (1 new gen., 2 new sp.) (mihirung birds).

¹ Those taxa prefaced by (x) were last studied by DeVis, and further study is needed to ascertain their correct taxonomic position.

- Northern Territory *Alcoota*. Waite Formation, Alcoota fauna (including Paine Quarry U.C.M.P. V-6345, Rochow Locality U.C.M.P. V-6349); cf. *Dromaius* (Casuariidae, emu), Dromornithidae (3 new gen., 3 new sp.) (mihirung bird), Accipitridae (eagle).
- New South Wales *Bugaldi*, diatomaceous carth deposit. cf. Aegothelidae (owlet-frogmouth).
- Victoria *Beamaris*. Sandringham Sands, Cheltenhamian Stage, Beaumaris fauna; Spheniscidae (penguin), ?*Pseudaptenodytes macraei* (Spheniscidae, penguin), ?*P. minor*, *Diomedea thyridata* (Diomedidae, albatross). Simpson, 1965, 1970; Wilkinson, 1969.
- Victoria *Spring Creek, Minliamite*. ?Cheltenhamian Stage; *Pseudaptenodytes macraei* (Spheniscidae, penguinidae).
- PLIOCENE
- South Australia *Lake Palankarina, Lawson-Daily Quarry (Lawson Quarry)*, Mampuworwu Sands, Palankarina fauna (U.C.M.P. V-5769); *dromaius ocybus* (Casuariidae, emu), Dromornithidae (mihirung bird). Miller, 1963a.
- New Guinea *Awe*. Otibanda Formation, Awe fauna; *Casuarius* sp. (Casuariidae, cassowar). Plane, 1967.
- South Australia *Lake Kanunka*. Katipiri Sands, Kanunka fauna (U.C.M.P. V-5772, V-5773, V-5855); *Dromaius* (Casuariidae, emu), *Pelecanus* (Pelecanidae, pelican), *Phalacrocorax* (Phalacrocoracidae, cormorant), (2 sp.), *Anhinga novaehollandiae* (Anhingidae, anhinga, snake bird), *Phoeniconaias gracilis* (Phoenicopteridae, flamingo), *Phoenicopterus ruber* (Phoenicopteridae) Anatidae (duck), cf. *Aythya* (Anatidae, duck). Stirton, Tedford, and Woodburne, 1967.
- Queensland *Peak Downs*. *Dromornis australis* (Dromornithidae, mihirung birds). Owen, 1874, 1879b.
- Queensland *Chinchilla*. Chinchilla Sands, Chinchilla fauna; *Dromaius* (Casuariidae, emu), *Pelecanus conspicillatus* (= *P. proavus*) (Pelecanidae, pelican), *Haliaeetus melanoleucos* (Phalacrocoracidae, cormorant), (x) *Xenorhynchus nanus* (Ciconiidae, stork), (x) *Anas elapsa* (= *Nettion elapsum*, Brodkorb)¹ (Anatidae, duck), (x) *Biziura exhumata* (Anatidae, duck), (x) *Dendrocygna* (Anatidae, duck), (x) *Nyroca reclusa* (= *reperta*, = *Aythya reclusa*, Brodkorb) (Anatidae, duck), (x) *Nyroca robusta* (= *Aythya robusta*, Brodkorb) (Anatidae, duck), *Gallinula* (= *Tribonyx mortierii reperta* (Rallidae, rail) (Olson, 1974), *Fulica atra* (Rallidae, rail) (Olson, 1974), DeVis, 1888-1905.
- PALEOGENE(?)
- Victoria *Redruth* (Wannon River). Ironstone, 'wing of a struthious bird', Aves, undetermined (David, 1950), feather impression (Chapman, 1910).
- MIDDLE TERTIARY
- Tasmania *Endurance Pit*. cf. Dromornithidae (mihirung birds), footprints. (Rich and Green, 1974).
- PLEISTOCENE
- QUEENSLAND
- Darling Downs *Dromaius* cf. *novaehollandiae* (Casuariidae, emu)
 Anhingidae (darter, J. van Tets, pers. comm., 1973)
 (x) *Palaeopelargus nobilis* (Ciconiidae, stork)
 (x) *Platalea ?subtenuis* (Threskiornithidae, spoonbill)
 (x) *Dendrocygna validipennis* (Anatidae, duck)
 (x) *Nyroca* sp. (Anatidae, duck)
 (x) *Necrastur alacer* (Accipitridae, hawk, eagle)
 (x) *Taphaetus brachialis* (= *Uroaetus* = *Aquila*) (Accipitridae, eagle)
Progura gallinacea (Megapodiidae, mound builder) (van Tets, 1974a)
Gallinula mortierii reperta (Rallidae, rail) (Olson, 1974) — ?
 Warwick
 (x) *Lithophaps ulnaris* (Columbidae, pigeon) DeVis, 1888-1905.

1. Generic reallocation most recently provided by Brodkorb, 1963, 1964, 1967, 1971a.

- Gore
Progura naracoortensis (Megapodiidae, mound builder). Longman, 1945; van Tets, 1974a.
- King's Creek, Darling Downs
 (x) *Dromaius patricus* (? = *D. novaehollandiae*) (Casuariidae, emu)
 (x) *Taphaetus brachialis* (= *Uroaetus* = *Aquila*) (Accipitridae, hawk, eagle). DeVis, 1888-1905.
- Diamantina (general locality; could be any of several places along the Diamantina River)
Genyornis (Dromornithidae, mihirung bird).
- Thorbindah, Paroo River
Dromaius (Casuariidae, emu)
Genyornis, cf. *newtoni* (Dromornithidae, mihirung bird).
- NEW SOUTH WALES
- Ashford Caves (Bone Cave) (U.C.M.P. V-67237, U.C.M.P. V-5545)
 Aves, undetermined.
- Bingara
Dromaius (Casuariidae, emu). L. Marcus, pers. comm., 1973.
- Canadian Lead
Dromornithidae (mihirung bird). Etheridge, 1889; Owen, 1879b.
- Cuddie Springs
Genyornis cf. *Newtoni* (Dromornithidae, mihirung bird). Anderson and Fletcher, 1934.
- Lake Menindee (including Rifle Range Loc. U.C.M.P. V-67233, U.C.M.P. V-5371, V-67185, V-67186, V-67187)
Dromaius (Casuariidae, emu).
- Lake Tandou
Dromaius (Casuariidae, emu)
Podiceps (Podicepedidae, grebe)
 cf. Ciconiiformes (herons, storks, ibises)
 cf. *Cygnus atratus* (Anatidae, swan)
 cf. Columbidae (pigeon).
- Wellington Caves (including Bone Cave, Mitchell's Cave, Phosphate Mine (in part U.C.M.P. V-5538))
 (x) *Dromaius patricus* (?*D. novaehollandiae*) (Casuariidae, emu)
Casuarius lydekkeri (Casuariidae, cassowary)
 Dromornithidae (mihirung bird)
Progura gallinacea (Megapodiidae, mound builder). Mitchell, 1839; Miller, 1962; Rich, 1973; van Tets, 1974a.
- Walli Caves
Progura gallinacea (Megapodiidae, mound builder). van Tets, 1974a.
- Wombeyan Cave (Guineacor) (in part U.C.M.P. V-5537)
 Aves, undetermined
Dromaius (Casuariidae, emu)
Progura gallinacea (Megapodiidae, mound builder). van Tets, 1974a.
- VICTORIA
- Buchan Cave (J. van Tets, pers. comm., 1973)
 Rallidae (rails)
Coturnix sp. (Phasianidae, quail)
 Alcedinidae (kingfishers)
 Passeriformes (song birds, perching birds).
- Lancefield
 Dromornithidae (mihirung bird).
- TASMANIA
- King Island
Dromaius minor (Casuariidae, emu). Spencer and Kershaw, 1910.

SOUTH AUSTRALIA

Baldina Creek (near Burra)

Dromornithidae (mihirung birds).

Brother's Island, Pt. Lincoln

Dromornithidae (mihirung bird).

Cooper's Creek (includes several localities collected primarily by J. W. Gregory and later parties from the University of California, Berkeley) (Katipiri Sands, Malkuni Fauna).

University of California Localities (Miller, 1963-1966; Rich, 1973)

- U.C.M.P. Site 2 (U.C.M.P. V-5379) Unkumilka Waterhole
Dromaius sp. (Casuariidae, emu)
Phalacrocorax sp. (Phalacrocoracidae, cormorant).
- U.C.M.P. Site 3 (U.C.M.P. V-5378) Between White Crossing and Site 2
Dromaius sp. (Casuariidae, emu)
Phalacrocorax sp. (Phalacrocoracidae, cormorant; middle-sized sp., lg. sp.).
- U.C.M.P. Site 4 (U.C.M.P. V-5380)
Anhinga novaehollandiae (Anhingidae, darter/snake bird)
Phalacrocorax sp. (Phalacrocoracidae, cormorant; middle-sized sp., lg. sp.)
Anatidae (duck).
- U.C.M.P. Site 5 (U.C.M.P. V-5381) Pirranna Soakage
Dromornithidae (mihirung birds)
Phalacrocorax (Phalacrocoracidae, cormorant; middle-sized sp.)
Anatidae (small sp.) (duck).
- U.C.M.P. Site 7 (U.C.M.P. V-5859)
Phalacrocorax sp. (Phalacrocoracidae, cormorant; middle-sized sp.)
Pelecanus conspicillatus (Pelecanidae, pelican)
Anatidae (duck)
Grus sp. (crane).
- U.C.M.P. Site 8 (U.C.M.P. V-5860)
Dromaius sp. (Casuariidae, emu)
Dromornithidae (mihirung birds)
Phalacrocorax sp. (lg. sp.)
Pelecanus conspicillatus (Pelecanidae, pelican)
Anatidae (swan)
- U.C.M.P. Site 9 (U.C.M.P. V-5861) Katipiri Waterhole (= Kuttipirra)
Dromaius sp. (Casuariidae, emu)
Dromornithidae (mihirung birds)
Phalacrocorax sp. (Phalacrocoracidae, cormorant, middle-sized sp.)
Anatidae (swan).
- U.C.M.P. Site 10 (U.C.M.P. V-5869)
Anhinga novaehollandiae (Anhingidae, darter/snake bird)
Phalacrocorax sp. (Phalacrocoracidae, cormorant)
Platalea sp. (Threskiornithidae, spoonbill).
- U.C.M.P. Site 14 (U.C.M.P. V-5866)
Anhinga novaehollandiae (Anhingidae, darter)
Phalacrocorax sp. (Phalacrocoracidae, cormorant)
Phoenicopterus ruber (Phoenicopteridae, flamingo).
- U.C.M.P. Site 16 (U.C.M.P. V-5868)
Phalacrocorax sp. (Phalacrocoracidae, cormorant)
Pelecanus conspicillatus (Pelecanidae, pelican)
Anatidae (swan).

U.C.M.P. Site 18 (U.C.M.P. V-6147)
Dromaius sp. (Casuariidae, emu)
Phalacrocorax sp. (Phalacrocoracidae, cormorant)
Pelecanus conspicillatus (Pelecanidae, pelican)
 Anatidae (duck).

Malkuni Waterhole (U.C.M.P. V-5382)
Pelecanus conspicillatus (Pelecanidae, pelican).

Markoni Locality (U.C.M.P. V-5382)
 Dromornithidae (mihirung birds)
Podiceps sp. (Podicepedidae, grebes)
Phalacrocorax sp. (Phalacrocoracidae lg. sp.)
Tyto (Tytonidae, barn owl).

J. W. Gregory Localities (DeVis 1888-1905; in Gregory, 1906)

East of Pirani

(x) *Biziura exhumata* (Anatidae, duck).

Emu Camp (= Malkuni waterhole)

cf. *Genyornis newtoni* (Dromornithidae, mihirung bird).

(x) *Phalacrocorax gregorii* (Phalacrocoracidae, cormorant)

(x) *P. vetustus*

(x) *Archaeocynus lacustris* (= *Cygnus lacustris*, Brodkorb) (Anatidae, swan)

(x) *Biziura exhumata* (Anatidae, duck)

(x) *Chenopsis nanus* (= *Cygnus lacustris*, Brodkorb) (Anatidae, swan).

Lower Cooper

Anhinga novaehollandiae (Anhingidae, darter)

(x) *Phalacrocorax gregorii* (Phalacrocoracidae, cormorant)

(x) *P. vetustus*

cf. *Dromaius novaehollandiae* (Casuariidae, emu)

(x) *Xenorhynchopsis tibialis* (Ciconiidae, stork)

(x) *Anas gracilipes* (= *Nettion gracilipes*, Brodkorb) (Anatidae, duck)

(x) *Archaeocynus lacustris* (= *Cygnus lacustris*, Brodkorb) (Anatidae, swan)

(x) *Chenopsis nanus* (= *Cygnus lacustris*, Brodkorb) (Anatidae, swan)

(x) *Nettapus eyrensis* (Anatidae, duck).

Lower Cooper Locality 2

Dromaius sp. (Casuariidae, emu)

Dromornithidae (mihirung birds).

Lower Cooper Locality 3

Dromaius sp. (Casuariidae, emu).

Lower Cooper Locality 4

(x) *Dromaius patricus* (cf. *D. novaehollandiae*) (Casuariidae, emu).

Lower Cooper Locality 5

(x) *Asturæetus furcillatus* (= *Plioæetus furcillatus* Richmond) (Falconidae, falcon).

Lower Cooper Locality 6

(x) *Baza gracilis* (= *Aviceda gracilis*, Brodkorb) (Falconidae, falcon).

Cooper Creek

Anhinga laticeps (Anhingidae, darter)

Pelecanus grandiceps (Pelecanidae, pelican).

Patteramordu (= Red Bluff Locality of U.C.M.P.)

(x) *Anas strenua* (= *Nettion strenum*, Brodkorb) (Anatidae, duck).

Unduwampa

- (x) *Xenorhynchopsis minor* (Ciconiidae, stork)
- (x) *Archaeocygnus lacustris* (= *Cygnus lacustris*, Brodkorb) (Anatidae, swan)
- (x) *Chenopsis nanus* (= *Cygnus lacustris*, Brodkorb) (Anatidae, swan).

Wankamamina

- (x) *Phalacrocorax gregorii* (Phalacrocoracidae, cormorant)
- (x) *Archaeocygnus lacustris* (= *Cygnus lacustris*, Brodkorb) (Anatidae, swan).

Wurdulumankula (Wurdulmankula)

- (x) *Dromaius patricus* (? cf. *D. novaehollandiae*) (Casuariidae, emu)
- (x) *Phalacrocorax gregorii* (Phalacrocoracidae, cormorant)
P. carbo (J. van Tets, pers. comm., 1973)
- (x) *P. vetustus*
- (x) *Xenorhynchopsis minor* (Ciconiidae, stock)
- (x) *Xenorhynchus nanus* (Ciconiidae)
- (x) *Ibis conditus* (= *Carphibis condita*, Brodkorb) (Threskiornithidae, ibis)
- (x) *Archaeocygnus lacustris* (= *Cygnus lacustris*, Brodkorb) (Anatidae, swan)
- (x) *Chenopsis nanus* (= *Cygnus lacustris*, Brodkorb) (Anatidae, swan)
- (x) *Nyroca effodiata* (= *Aythya effodiata*, Brodkorb) (Anatidae, duck)
- (x) *Leucosarcia proevisa* (Columbidae, pigeon).

Henschke's Quarry Cave, Naracoorte

- Progura naracoortensis* (Megapodiidae, mound builder). van Tets, 1974a.

Kangaroo Island

- Dromaius novaehollandiae diemenianus* (Casuariidae, emu). (Condon, 1973).

Lake Callabonna (Lower level)

- Dromaius* sp. (Casuariidae, emu)
- Genyornis newtoni* (Dromornithidae, mihirung bird).

Lake Callabonna (Upper level)

- Phalacrocorax* sp. (Phalacrocoracidae, cormorants).

Lake Eyre (exact locality unknown)

- (x) *Ocyplanus proeses* (Laridae, gull). DeVis in Gregory, 1906.

Mount Gambier Range

- Dromornithidae (mihirung birds).

Normanville (= Salt Creek)

- Dromornithidae, cf. *Genyornis newtoni* (mihirung bird).

Penola

- Dromornithidae, cf. *Genyornis newtoni* (mihirung bird).

Victoria Cave, Naracoorte (van Tets and Smith, 1974)

- Dromaius* sp. (Casuariidae, emu)
- Progura naracoortensis* (Megapodiidae, mound builder)
- Leipoa ocellata* (Megapodiidae, mound builder)
- Coturnix pectoralis* (Phasianidae, quail)
- C. australis*
- Coturnix* sp. (Phasianidae, quail)
- Turnix varia* (Turnicidae, button quail)
- Pedionomus torquatus* (Pedionamidae, plains wander)
- Rallus philippensis* (Rallidae, rail)
- Peltohyas australis* (Charadriidae, plovers)
- Tringa glareola* (Scolopacidae, sandpiper, stilt, curlew)
- Gallinago hardwickii* (Scolopacidae, sandpiper, stilt, curlew)

Calidris ruficollis (Scolopacidae, sandpiper, stilt, curlew)
Chionis minor (Chionidae, sheath-bill)
Pezoporus wallicens (Psittacidae, parrot)
Tyto novaezelandiae (Tytonidae, barn owl)
Gymnorhina tibicen (Passeriformes, song birds).
Gymnorhina tibicen (Passeriformes, song birds)

Warburton River Localities (Katipiri Sands, Mulkuni Fauna)

Cassidy Locality (U.C.M.P. V-5539)
 cf. *Genyornis* (Dromornithidae, mihirung bird).

Green Bluff Locality (U.C.M.P. V-5775)

Dromains sp. (Casuariidae, emu)
Phalacrocorax (Phalacrocoracidae, cormorant; lg. sp.)
 Anserinae (goose)
 Accipitridae (hawk, eagle).

Kalamurina

Dromains sp. (Casuariidae, emu)
 Dromornithidae, cf. *Genyornis* (mihirung bird)
 (x) *Phalacrocorax gregorii* (Phalacrocoracidae, cormorant)
 (x) *P. vetustus*
 (x) *Anas gracilipes* (= *Nettion gracilipes*, Brodtkorb) (Anatidae, duck)
 (x) *Taphaeus lacertosus* (= *Icthyophaga lacertosus* (van Tets, 1974c)
 (Accipitridae, eagle)). DeVis, 1888-1905; Rich, 1973.

Lookout Locality (U.C.M.P. V-5776)

Phalacrocorax sp. (middle-sized sp., lg. sp.) (Phalacrocoracidae, cormorant).

Marcus Locality

Phalacrocorax sp. (Phalacrocoracidae, cormorant; middle-sized sp., lg. sp.)
 cf. *Pelecanus* sp. (Pelecanidae, pelican).

Pundrakadarinna Soakage (U.C.M.P. V-5777)

Phalacrocorax sp. (Phalacrocoracidae, cormorant).

Warburton River

(x) *Pelecanus validipes* (Pelecanidae, pelican). Etheridge, 1894.

Weeke's Cave (Nullarbor Plain)

Platibus flavipes (Threskiornithidae, spoonbill)
Accipiter fasciatus (Accipitridae, hawks)
Falco cuculoides (Falconidae, falcon)
Turnix sp. (Turnicidae, bustard quail)
Cincloramphus cruralis, *C. mathewsi*, *Poephila guttata*, *Artamus leucorhynchus* (Passeriformes, song birds). van Tets, 1974b.

WESTERN AUSTRALIA

Labrinth Cave

Aves, undetermined.

Madura Cave

Aves, undetermined.

Mammoth Cave

Dromornithidae (mihirung bird).

Scott River

?Dromornithidae (mihirung bird), egg. Butler, 1969; Rich, 1973.

Previous Opinions Regarding Origin of the Australian Avifauna

In studying the Australo-Papuan avifauna three decades ago, Mayr (1941, 1944a-b, 1945b) concluded that it had been derived from the north, primarily southeastern Asia, with colonization occurring throughout the Cenozoic (see figure 9a). He recognized several groups of invaders, which he believed to represent different colonizations, oldest to most recent. The oldest invasion (early to mid-Tertiary at the latest, Mayr, 1944a) was reflected by the most unique groups in the living non-passeriform avifauna of Australia, including the Casuariidae (containing *Casuarinus* and *Dromaius*), Megapodiidae, Loriinae, Caeatuinae (= Kakatoeinae), Platyeereinae, and Podargidae. Mayr noted that the nearest relatives of these six groups were uncertain, but argued that none were any more closely related to South American families than to Old World families. His second and more recent wave of colonists included the Pedionomidae (collared hemipodes) along with several passeriform families. The third wave consisted of a number of genera endemic to Australo-Papua, which he believed were Mioene or Pliocene arrivals, and included: *Synoicus* (Phasianidae), *Geopelia* (Columbidae), *Irediparra* (included in *Jacana*, Jacanidae), *Notophoxyx* (included in *Ardea*, Ardeidae), *Dupetor* (Ardeidae), *Erythrotriorchis* (Accipitridae), *Uroaetus* (= *Aquila*, Accipitridae), *Lophoictinia* (Milvinae), *Syna* (included in *Halcyon*, Alcedinidae), *Dacelo* (Alcedinidae), *Tanysiptera* (Alcedinidae), *Cacomantis* (Cuculidae), *Misocalius* (included in *Chrysococcyx*, Cuculidae), *Chaleites* (included in *Chrysococcyx*, Cuculidae), and *Eudynamis* (Cuculidae). Fourth and fifth waves included taxa that are presently only specifically, or subspecifically distinct, or indistinguishable from southeast Asian forms (Pliocene to Recent arrivals, according to Mayr). Such ideas viewed against a backdrop of stable continents were reasonable and remained so into the late 1950's. With the development during the last fifteen years of the concepts of continental drift, sea-floor spreading, and plate tectonics, it has become apparent that two

major routes may have existed that non-marine vertebrates could have employed in reaching or leaving Australia during the late Mesozoic and Cenozoic: the Antarctic as well as the Indomalaysian. With this additional information provided by geology, one of Mayr's basic assumptions, that of continental stability, is undermined, and his ideas regarding the origin of the Australian avifauna require re-examination.

Such re-evaluation has been initiated in papers by Serventy (1972, 1973), Cracraft (1972, 1973), Schodde and Calaby (1972), and Mayr (1972) himself. Serventy suggested such forms as *Dromaius* (emus), possibly *Anseranas* (Magpie Goose), the Megapodiidae (mound builders), the Podargidae (frogmouths), as well as several parrot subfamilies (Loriinae, Caeatuinae, Platyeereinae (included in the Psittacinae)) among the non-passeriforms might well have utilized a southern, rather than a northern route between Australia and the remaining world. Mayr, in commenting on Serventy's paper, tentatively considered the ratites, the platyeereine parrots, and possibly the flamingos as Antarctic dispersers but left the question open. Cracraft, on the other hand, strongly supported a southern dispersal route for the ratites, megapodes, and parrots, but agreed with both Serventy and Mayr that the majority of Australian avian genera (including most of the non-passeriforms) were spawned by Southeast Asian faunas. Is this really the case, however, and what sort of reasoning and assumptions are employed to determine each family's route of dispersal between Australia and the rest of the world? The following section deals in detail with these questions.

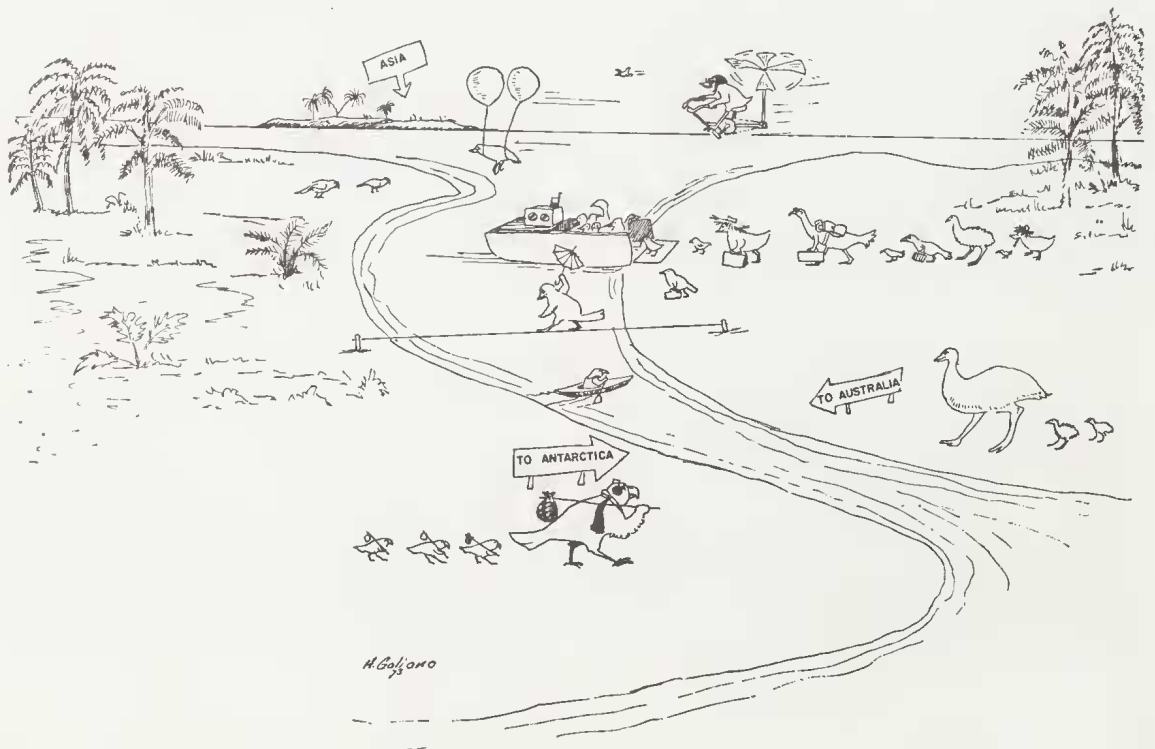
Re-evaluation of Probable Immigration Routes Into Australia: Assumptions, Reasoning, and Data

Zoogeography is often not a very satisfying endeavour owing to the frequent uncertainty of conclusions one is able to reach concerning areas of origin and early radiation of certain organisms. Despite this, however, many workers have attempted to understand the distributional histories of organisms including birds, although



Fig. 9—(A) Cartoon expressing the classic ideas concerning origin of the Australian avifauna, eloquently detailed by Mayr (1944a-b); (B) cartoon depicting an additional component

that together with Mayr's ideas may have been important in shaping the composition of Australia's bird fauna.



others seriously question many of the criteria utilized in such attempts (Rotramel, 1973, among others). The following section summarizes briefly assumptions and reasoning utilized by ornithologists and zoogeographers who have tried to determine origins of elements comprising the Australian avifauna. In addition, it outlines assumptions employed in this paper to re-evaluate the immigration and emigration pathways of all the non-passeriform families, presents pertinent historical, distributional, and diversity data for each family considered, and briefly outlines the reasoning employed in the determination of the dispersal route(s) utilized by each Australian group between Australia and the remaining world.

In his treatise on zoogeography, Darlington (1957) outlined several working principles and a number of practical clues used in determining the area of origin of different organisms including (1) diversity, (2) degree of differentiation (including degree of endemism), (3) extent of area inhabited (whether restricted or widespread), (4) continuity of area inhabited, (5) distribution of related, competing, and associated families, and (6) fossil record. How much weight to give each of these clues, particularly in cases where information is limited, was not made clear. Similarly, if some data are available for each clue category, it was not made clear which clue or clues had priority over others if multiple interpretations were possible. Darlington suggested that degree of differentiation (clue 2) should have priority over diversity (clue 1), in fact that information pertinent to the remaining clues (clues 3, 4, 5, 6) could be used to question the significance of diversity. He quite appropriately pointed out that high diversity in an area might not only signal the long history of a group there, but might also be due to a number of other factors such as the size and diversity of the environment inhabited (i.e. number of different habitats available), and the amount of competition a group encountered from other organisms during its history in an area. To this list Keast (1972) added such factors affecting group diversity as size of total area inhabited, topographic diversity, percentage of area within the tropics, total latitudinal range on continent, vegetational

composition of continental flora, and history of connection to other land masses. Darlington further stressed other problems peculiar to deciphering avian zoogeography including the (1) rapidity with which an avian taxon can disperse, thus obscuring the direction from which it came; (2) possibility of contemporaneous multiple dispersals; (3) rarity of avian fossils; and (4) lack of understanding of the significance of non-breeding ranges of migratory birds. However, after so carefully and precisely outlining these assumptions and clues, most were not evaluated when Darlington individually considered a number of "dominant" avian families; instead diversity and endemism were usually the only clues considered. Darlington's judgement on the relative significance of these few clues was not consistent; on the one hand he believed that the Phasianidae originated, or at least underwent a major radiation, in the Oriental region based on their great diversity there⁶ (in comparison to its low diversity elsewhere), but was unwilling to readily accept the equal probability that the parrots originated or underwent an early radiation in Australia based on the same type of distributional and diversity information.⁷ Besides pheasants and

⁶ "It [the Phasianidae] is the first large family of this survey which shows an apparent pattern of radiation from one main center, the Oriental Region. The true pheasants, jungle fowls, and peacocks are almost all in the Orient, and from there the ancestors of *Afropavo* (the Congo Peacock), and of the guinea fowls may have reached Africa; the ancestors of the grouse (Tetraoninae) may have gone northward and around the northern part of the world, eventually differentiating most in North America; and the quails may have radiated over the whole world, with a secondary evolution center in southern North America. The fowl-like birds *probably* [italics mine] have relatively low powers of dispersal, so their present distribution may reflect something of their early history . . ." Darlington (1957, p. 270).

⁷ "Parrots . . . are almost equally numerous in species in Old and New Worlds but are much more diverse in the former, all the New World forms belonging to one of the several Old World subfamilies; this suggests an Old World origin of parrots. But this does not carry the analysis far enough. It is only in the Australian Region and adjacent island areas that parrots occur in great diversity. Elsewhere in the Old World parrots are relatively few and all belong to one subfamily, the same one that is in America. Does the unique diversity of parrots in the

parrots, Darlington examined several other avian groups that occur in Australia (Anatidae, Falconiformes, Strigiformes, Ardeidae, Charadriidae, Scolopacidae, Rallidae, Cuculidae, Caprimulgiformes, Apodidae, Alcedinidae), attempting to determine their centres of origin and distributional histories, but could reach no further firm conclusions for any other non-passeriform group. He believed that most of the groups which he considered were old, had complex distributional histories, and that areas of origin as well as early patterns of radiation could not be determined. He noted that the Caprimulgiformes (nightjars, frogmouths, and allies) probably had experienced three major radiations, one in the Australian-Oriental area (Podargidae, Aegothelidae), another in tropical America (Nyctibiidae, Steatornithidae), and a third in an undesignated area that produced the Caprimulgidae, but he was non-committal about what dispersal paths had been employed during this tripartite diversification. It is with this background of uncertainty about the dispersal histories of birds that Darlington made the following statement: "Almost all the ascertainable relationships of Australian land birds are toward Asia. The few supposed African relationships, for example that of the honey eaters with *Promerops* of South Africa, are doubtful, and there is no sure, direct relationship between any Australian and South American land birds" (1957, p. 262). Thus with this statement and others regarding the origins of the parrots and pigeons, Darlington emphasized the "Asiatic" character of the Australian avifauna, essentially agreeing with Mayr's ideas.

Mayr's (1944a-b) reasoning, that most Australian birds were derived from Oriental forms, relied heavily on the two observational assumptions that (1) the Australian avifauna more closely resembled that of the Oriental region than that of any other area and (2) during the history of birds, Australia had re-

mained in its present geographic position, with colonization slowly accelerating as more of the Malay Archipelago emerged during the Cenozoic. With such a foundation, Mayr's conclusion (that Australia's avifauna consisted of several different "waves" of northern colonists) was quite reasonable. However, as a previous section of this paper has established, present geological evidence strongly suggests that Australia has been anything but stable during the history of birds, having drifted as much as 30-50° of latitude from an Eocene connection with Antarctica nearly to its present position by the late Miocene. Thus Mayr's second assumption is seriously challenged. If his first assumption is re-examined, although most certainly true, it could have resulted from at least two types of faunal interchange, i.e. (1) Australia receiving most of its avifauna from Asia or (2) the most recent biotic interchange between Australia and the rest of the world being with Asia. If the latter is the case, similarities between Asia and Australia would have resulted from a mixing of the two faunas, not a one way immigration from Asia. Because of the present lack of understanding of avian distributional histories, however, the conclusion that Asia spawned most of Australia's avifauna seems premature, a conclusion far too specific for the limited data available.

Serventy (1972, 1973), accepted the recent consensus of geologists supporting continental drift and re-examined the old, endemic elements in Australia's avifauna, attempting to point out the closer relationships of some Australian birds to South American rather than Asian forms. His acceptance of ratite monophyly (as most recently supported by Bock, 1963; Meise, 1963; Parks and Clark, 1966; Sibley, 1960) allowed him to support strongly a southern dispersal route for this group, and along with additional data, led him to state: "May not some of the older elements of the present Australasian fauna have reached Australia via Africa and Antarctica during the earlier stages of Gondwanaland?" Serventy did not extend his considerations to other, less unique Australian bird groups, because he believed they had immigrated from Asia due to their lack of close similarity to South American forms.

Australian Region reflect place of origin, or withdrawal of some groups of parrots from other parts of the world, or local radiation? I cannot answer this question, but it seems to me that parrots need not have originated in the Australian Region . . ." Darlington (1957, p. 271).

Cracraft (1972, 1973) likewise assumed that continental drift had occurred during the Mesozoic and Cenozoic (in fact he presented a review of continental drift literature) and attempted to elucidate the dispersal histories of avian taxa based primarily on phyletic relationships of those taxa (using a method discussed by Hennig, 1966; Nelson, 1969; and others). He emphasized that all other clues regarding origin of avian groups should be subordinate to phyletic clues, but then noted how pitifully poor present understanding of phyletic relationships was for most avian families and thus, I assume, how little is understood of original dispersal routes if that information is derived from phyletic studies. He discussed those few groups whose relationships he believed were best understood, including taxa with Australian histories: penguins (Spheniscidae), ratites, galliforms (including Megapodiidae), parrots (Psittacidae), pigeons (Columbidae), and cuckoos and allies (Cuculiformes) among the non-passeriforms. Yet with this limited consideration as a basis, he was willing to conclude in another paper (Cracraft, 1972, p. 173, as well as 1973) that "the vast majority of Australian families undoubtedly have relationships to Asian families and therefore can be supposed to have come from the north. This includes *most non-passerines* [*italics mine*] and all oscines." Cracraft, like Serventy, did not present data that would support such a statement, and thus presumably accepted much of the reasoning previously articulated by Mayr and Darlington. Unfortunately, all of these workers have not stated the important point that similarity of the Australian and Asian avifaunas could well be due to a two-way, not just a one-way, exchange of old endemic faunas, and that present-day avifaunal similarity need only signal the latest, not necessarily the sole, episode of interchange between Australia and the rest of the world. Why not investigate the Australian nature of the Oriental avifauna as well as the Asian nature of the Australian avifauna? It, also, should be realized that because linkage between Australia and other southern continents was broken during or before the Eocene, birds on any of these gondwanic continents today have had ample

time to evolve quite separately and diverge from one another. Relationships between living forms that might be used in support of such an early Tertiary dispersal route should thus be sought at higher taxonomic levels, not specific and generic.

In this paper a number of assumptions will be made and clues used in an attempt to decipher the dispersal history of Australian avian families. Assumptions include:

- (1) the geographic arrangement of continents has been continually altered throughout the history of birds and quite profoundly in the positioning of Australia during the Cenozoic (i.e. Australia has drifted from an early Tertiary position adjacent to Antarctica, perhaps as much as 30-50° of latitude northwards), nearing its present position by the late Miocene;
- (2) if the geographic position of a continent relative to others has changed during the history of a group of organisms, then faunal composition on any one continental mass may reflect several episodes of exchange with other geographic areas; those areas most recently in contact or closely apposed to one another should be expected to exhibit the greatest faunal similarity.

With these assumptions the practical clues examined in an attempt to determine the dispersal routes to and from Australia utilized during the history of any one avian family include:

- (1) phyletic relationships of the family to others (accompanied by distributional and degree of differentiation data for "other" families);
- (2) fossil history of family, worldwide;
- (3) degree of endemism of family both within and outside of Australia (see figure 5);
- (4) diversity of family both within and outside of Australia (see figure 5);
- (5) nature of distribution: cosmopolitan, pantropical, restricted; continuous or segmented.

How each of the above elucs are interpreted in the following study needs brief explanation:

Phyletic relationships. If a group (or groups) can be identified as being closely related to a family with Australian representatives, and information is available concerning intragroup phyletic relationships, then those data will sometimes favour one dispersal route over another between Australia and the rest of the world, although not always allowing resolution of the direction of movement along that route. An explanation of the observed distribution of the group members (including the Australian taxa) attempts to determine the minimum number of dispersals necessary (see Nelson, 1969, for excellent discussion of this procedure) to produce the present array of birds with the constraints imposed by understanding of phylogenetic relationships and knowledge of Mesozoic and Cenozoic paleogeography. Such explanations, however, offer only "parsimonious" first approximations that may well be refined, and perhaps found to be decidedly more complex, if understanding of the phylogeny of certain avian groups is improved and/or a more geographically and temporally complete fossil record of birds materializes in the future. Recent and fossil material are of equal importance when constructing phylogenies, each limited by incompleteness of specimens (or study thereof) or of geographic representation. Geographic position of the most primitive taxa (especially if based on living forms alone) is *not a priori* assumed to be indicative of the centre of origin or early radiation of a group but must be examined in light of the distributional patterns of other more primitive and advanced members of the group.

Unfortunately, this clue is of limited use because of the poor understanding of phylogenetic interrelationships of many of the non-passeriform taxa. This is well demonstrated by diversity of opinions expressed today concerning interfamilial relationships of birds, many of which are cited in the following section.

Degree of endemism and diversity. Darlington (1957) has posed some serious objections to the use of endemism and diversity (see previous section of this paper), but in this study, such clues do seem helpful in two

extreme cases, i.e. where (1) a family is highly diverse, having a large number of endemic genera/subfamilies in Australia but is unrepresented or has only a scant record elsewhere and (2) a family is highly diverse outside Australia but is represented in Australia by a single genus or species, often conspecific with extra-Australian populations.

Case 1 may represent a group that has had a long history in Australia and just recently emigrated elsewhere, especially if the extra-Australian forms are closely (conspecific or at the most congeneric) related to Australian taxa. The high diversity and endemism in Australia cannot be explained by that continent's greater environmental diversity in comparison to that of its nearest continental neighbour today, the Palearctic-Oriental region. Certainly a greater topographic and climatic diversity, a decidedly greater land area (ca. 3 million miles² in Australia; ca. 21 million miles² in the Oriental-Palearctic; and ca. 33 million miles² in the Old World when Africa is included (Golenpaul, 1960)), as well as a greater tropical land area characterizes the Palearctic-Oriental Realms when compared with the Australian (excluding New Zealand). As previously mentioned (also see below), the effect of differential competition encountered by an avian group in either area cannot be measured and remains unknown in this consideration. Thus original entry of groups in this category into Australia via an Antarctic dispersal route should be considered a possibility along with an Indomalaysian route. If the extra-Australian forms are quite distinct (although few in number), however, the situation may be more complex and will be discussed further under each family affected.

Case 2 apparently represents the opposite extreme, a group that has only recently arrived in Australia and has not yet differentiated to any marked degree. In this case the Indomalaysian or transoceanic route is most probable paths of original entry.

Of course, these two cases represent the extremes of a spectrum of possibilities; many avian families are moderately diverse, with many endemic taxa in several areas of the world, including Australia. Unless other data

are available, it may be impossible, as Darlington (1957) has suggested, to determine the most probable route of initial entrance for these families at present.

Nature of distribution. Distributional clues have been used as discussed by Darlington (1957), i.e. the continuity or discontinuity of an area inhabited by a particular group can be used to determine if a group has recently or long ago invaded an area, where a disjunct distribution is indicative of a group that has invaded long enough in the past to have experienced fragmentation into disjunct populations separated by considerable distances.

Two other clues that Darlington (1957) employed in his considerations of avian zoogeography are not particularly useful when considering the origin of the Australian avifauna because of the total lack of or rarity of data for each: the distribution of "competing" families and the composition of fossil avifaunas.

In the former case I know of no careful and sufficiently broad-based studies designed to determine what avian families (or for that matter what other organisms, not just birds) are "competing" with certain Australian taxa both within and outside of Australia. How much (or how little) competition a taxon encounters in different geographic areas may well be the primary factor accounting for differential diversity in certain areas, but how can this be recognized and measured? Darlington (1957) very appropriately pointed this problem out, and it should be kept in mind to temper all conclusions regarding location of early centres of radiation, as well as antiquity of families in certain areas, particularly in the case of Australia where the early Tertiary fauna may well have been impoverished and unbalanced and where there is no fossil record to confirm what groups were actually there.

The fossil record, also, is not particularly helpful at present in determining what dispersal route was used by birds present in Australia during the Cenozoic because of the incompleteness of the Paleogene record in Australia (see Table 3), South America (Pascual and Rivas, 1971), and Southeast Asia. Only Europe and to a lesser extent North America (mid to

late Cenozoic in particular) have produced sufficient fossil material to allow partial reconstruction of Cenozoic avifaunas. To some extent, early Tertiary European avifaunas can be used to suggest what avian taxa might have simultaneously lived in Southeast Asia, but such would only be extrapolation in need of confirmation, since barriers to overland dispersal may well have separated the two areas during parts of the Tertiary (see Szalay and McKenna, 1971; McKenna, 1972a). Just because such early Tertiary faunas are not presently known, however, one should not condemn the value of fossils for the future. Were the records more complete, particularly in South America and Australia, questions regarding the antiquity of many groups in these areas could be answered, as could those concerning which dispersal route between Australia and the rest of the world was most likely for those groups. Were the record more complete, it could be used to evaluate parsimonious conclusions based solely on phyletic relationships and distribution of modern taxa, which may be gross oversimplifications of reality.⁸ In making such a statement, I am not advocating a cessation of research on avian phylogeny, which I think can

⁸ A group that well exemplifies such oversimplification by parsimonious arguments based on the living fauna alone is the mammalian order Perissodactyla represented by the tapirs (tropical America, Southeast Asia, and Indonesia), the rhinoceroses (Africa, eastern Asia, and Indonesia), and the horses and allies (central and eastern Asia and Africa) in the modern fauna (Walker, 1964). Of these living forms the tapirs are the most primitive and the horses most advanced or derived (Simpson, 1945). Thus the most parsimonious explanation of origin and dispersal of this group would require an Old World origin of the equids and the rhinoceroses and an Old or New World origin of the tapirs. The fossil record, however, indicates the "reality" of perissodactyl dispersal was far more complex than the modern record intimates. A wide variety of equids and rhinoceroses as well as a number of now extinct perissodactyl groups inhabited North America and Europe, as well as those areas now inhabited by this order at various times during the Cenozoic. Without necessarily altering the hypothesized phyletic relationships of the living perissodactyls, the fossil evidence certainly increases the possibilities for area of origin and early radiation of both the horses and rhinoceroses, challenging the specificity implied by the original parsimonious argument.

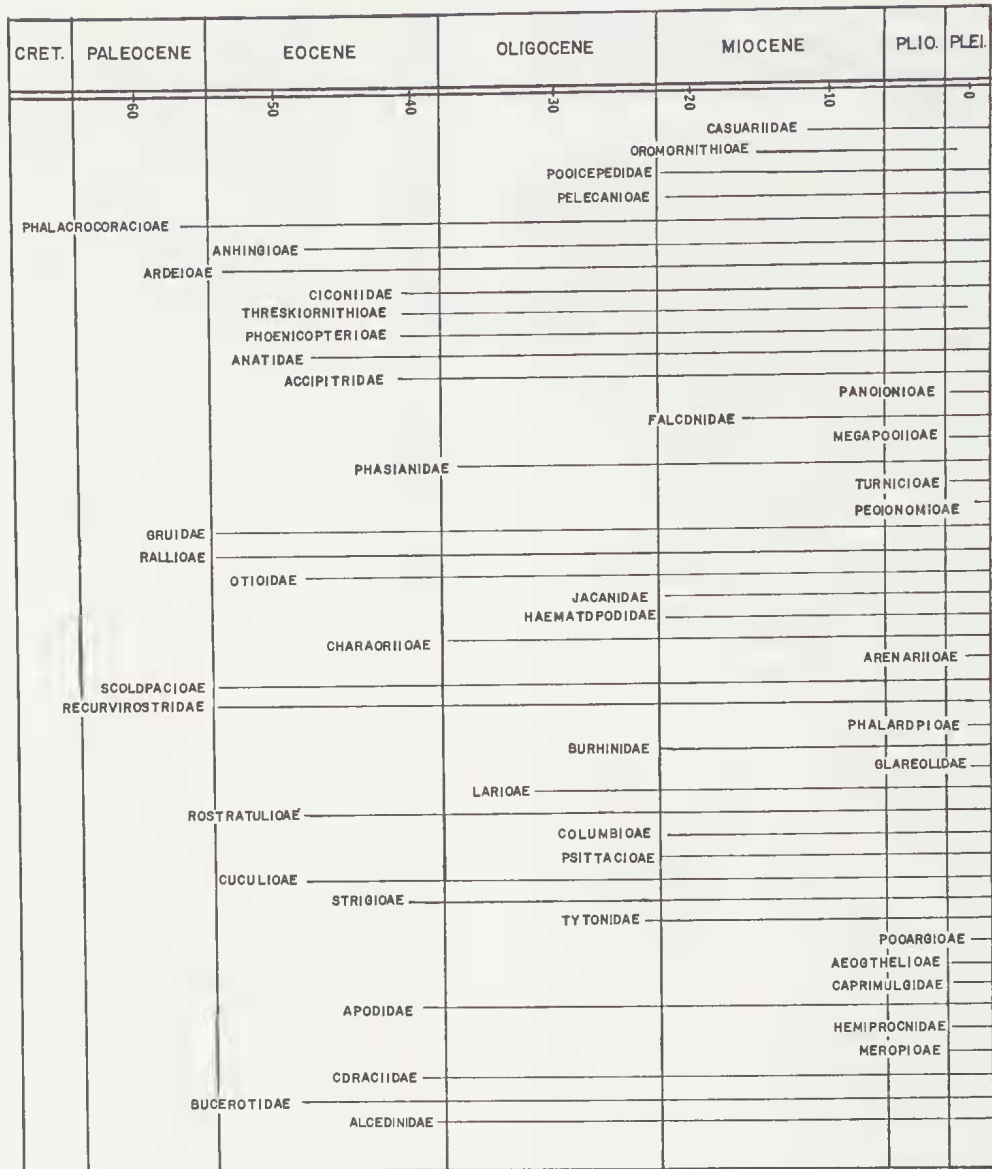


Fig. 10—Geologic ranges (world-wide) for those non-passeriform avian families represented in the Cenozoic of Australia. Horizontal axis calibrated in millions of years, in general following Berggren, 1969, 1971.

only help in understanding of avian distribution, but I am criticizing those who would condemn the fossil record as a useless, even misleading item of information for zoogeographic considerations. A good fossil record combined with data from the living fauna and continental reconstructions for the Mesozoic and Cenozoic, on the contrary, can offer alternatives (and often a closer estimate of reality, even if that

estimate is more complex) that cannot be confirmed by, or deduced from, the modern avifauna alone (see Cracraft and Rich, 1972, concerning the early radiation of the Cathartidae in Europe; also Nelson, 1969).

Despite the rarity of avian fossils in most parts of the world, the palaeontological record is important to this study in one way. It demonstrates that at least twenty of the forty-six avian families (see figure 10) that make up the Cenozoic avifauna of Australia had probably differentiated by the Eocene (data mainly from Brodkorb, 1963, 1964, 1967, 1971a) and thus

could have utilized the Antarctic dispersal route to initially disperse between Australia and the remaining world.

Summary of Phyletic Relationships, Diversity, Endemism, Distribution, and Palaeontological Record of Avian Families Having Australian Representatives

Using the five clues (phyletic relationships, diversity, endemism, distribution, fossil record) stressed in the previous section, an attempt is made in this section to determine for each avian family with a Cenozoic record in Australia what the most probable route of entry has been. Palaeontological data are given to emphasize which families are known to have differentiated by the early Tertiary, thus indicating the real possibility that the Antarctic dispersal route could have been used. Data for each family are presented in the following order: technical and common names; temporal range (both for world and Australia; if coincident only one range listed); geographic range; total number of living and fossil genera (if different from following category); total number of living and fossil genera in Australia-New Guinea; geographic area of greatest diversity; geographic area of greatest endemism (at generic, sub-familial levels); phyletic relationships to other avian families. A brief analysis follows each set of data for every family and attempts to establish which dispersal route, Antarctic and/or Indomalaysian, has been used. In many cases both routes may have been utilized at different times during the history of an avian family while in other cases one route seems more probable. In several cases, a choice of routes cannot be made based on available data.

The familial classification employed follows that of Peters (1931, 1934, 1937, 1940, 1945) with synonymies of the Mayr and Amadon (1951) and the Brodkorb (1963, 1964, 1967, 1971a) classifications also listed. Generic assignment to each family follows Peters (see above) except where modified by Bock (1956), Woolfenden (1961), Brown and Amadon (1968), Rand and Gilliard (1968), Sibley and Frelin (1972), Rich (1973) and Condon (in press). Data on diversity and endemism were derived primarily from Peters and Condon

(see above), palaeontological data from Brodkorb (see above and 1971b) and Olson (1974), and phylogenetic information from Brodkorb, Mayr and Amadon, Storer (1960, 1971), Woolfenden, Brown and Amadon, Bock, and Sibley and Ahlquist (1972). The latter reference summarizes most of the important works on avian phylogenetic studies and will not be duplicated here.

Casuariidae (Dromiceidae, Casuariidae of Mayr and Amadon, 1951; Brodkorb, 1963; Casuariidae of Sibley and Frelin, 1972). Emus and cassowaries; Miocene-Recent; *Dromaius* restricted to Australia, *Casuarius* to Australia, New Guinea, and some Pacific islands; two genera in family, *Dromaius* and *Casuarius*; greatest endemism and diversity in Australia; nearest extra-Australian relatives, elephant birds and rheas-ostriches, not moas and kiwis; nearest relatives, the dromornithids (Rich, 1973).

The emus and cassowaries form a unique, quite distinct and moderately diverse ratite group in Australia whose nearest extra-Australian (New Guinea and Southwest Pacific included) relatives occur in South America and Africa (see Rich, 1973; Cracraft, 1973; Sibley and Frelin, 1972; Jehl, 1971; Glenny, 1965; Sibley, 1960). The family is also present, but neither as diverse nor endemic, in New Guinea and on some southwest Pacific islands. The only ratite known from Asia, the ostrich (*Struthio*), first recognized in Europe and Africa in the late Miocene (Rich, 1972), is far too specialized to have given rise to the Australian ratites, however. Casuariid differentiation had occurred in Australia by the Miocene, and therefore a Paleogene or earlier differentiation is necessitated. More advanced ratites were present in South America during the Eocene (Brodkorb, 1963; Rich, 1973; Cracraft, 1973), and thus forms that could have been ancestral to Australian ratites must have differentiated even before, and an early dispersal of this group across a southern route must be considered a possibility.

Dromornithidae. Mhirung birds; Miocene-Pleistocene (presently extinct); restricted to Australia; 5-6 genera, including *Genyornis* (Stirling, 1896; Stirling and Zietz, 1896; 1900;

1905; 1913), *Dromornis* (Owen, 1872; 1874; 1879a-b) and 3 new genera (Rich, 1973); nearest extra-Australian relatives probably elephant birds (Acipornithidae), rheas (Rheidae); nearest relatives the Casuariidae.

The Mhirung Birds (Rich, 1973) or "giant emus" appear to be most closely related to the Casuariidae (Rich, 1973) and both in turn are closer morphologically to the more primitive elephant birds and more advanced rheas-ostriches than to the moas-kiwis (Rich, 1973). The same biogeographic arguments that are valid for the casuariids hold for the Dromornithidae, and thus an Antarctic route needs to be considered a possibility for the ancestors of both groups.

Podicipedidae. Grebes: Miocene-Recent; Pleistocene-Recent in Australia; cosmopolitan distribution; 6 genera in family (2 extinct); 1-3 genera (*Podiceps*, ?*Poliocephalus*, ?*Tachybaptus*) having cosmopolitan distribution, with remaining genera in New World; nearest extra-Australian relatives (at the family level) not certain, perhaps the loons (Gaviidae) with an entirely Holarctic distribution (first fossil occurrences Paleocene of Europe, Oligocene of North America), Enaliornithidae (extinct, Cretaceous of England), Lonchodytidae (extinct, Cretaceous of North America), Baptonithidae (extinct, late Cretaceous of North America), Charadriiformes, Spheniscidae, Pelecaniformes, Procellariiformes, or Anseriformes.

Three species of grebes (*Podiceps ruficollis*, *P. cristatus*, and *P. poliocephalus*) are present in the recent Australian avifauna. The first two have ranges extending into most of the Old World, and only *P. poliocephalus* is endemic to Australia. For a number of reasons initial entrance via southward dispersal along the Indomalaysian route seems most probable: (1) greatest generic diversity within the family as well as specific diversity within *Podiceps* is in the New World; (2) probable nearest relatives to the family occur in Holarctica (including North America); and (3) the family is represented by only one genus in Australia, a genus that is not endemic or particularly diverse there.

Pelecanidae. Pelicans: Miocene-Recent; one genus (*Pelecanus*) with world-wide distribution

and a second fossil genus, *Liptornis* (mid-Miocene, South America); nearest extra-Australian relatives probably within the Pelecaniformes: Lanham (1947) believed the Phaethontidae and Fregatidae, with recent pantropical distributions, were the most primitive members of the Pelecaniformes, and thus more primitive but closely related to the Pelecanidae; other possible close relatives are the fossil families Odontopterygidae (Eocene of England), Pseudontornithidae (Miocene of Brazil or Germany?, Miocene of North America, Pliocene of New Zealand), Pelargornithidae (Miocene of Europe), Cladornithidae (Oligocene of South America), Cyphornithidae (Miocene of North America).

More diverse in the past (2 genera in the Miocene), the pelicans are represented by a single genus in the living avifauna that is most varied in the Old World (Africa-Europe-Asia). One species occurs in, and is endemic to, Australia, as is another in the New World (out of a total of seven species). Because of the lack of distinct endemics (at the generic level) restricted to Australia and/or Australia-Old World and the low diversity of the genus *Pelecanus* in Australia, entrance via the Indomalaysian route or even oceanic dispersal is more probable. Also, if the Pelecanidae really were derived from forms similar to tropic birds and frigate birds with tropical distributions, Indomalaysian dispersal would be favoured.

Pelecanus tirarensis, the mid-Miocene form from interior Australia, may well be close to the ancestral form that first entered Australia (as it neared Southeast Asia) and gave rise to the living *P. conspicillatus*.

Phalacrocoracidae. (Phalacrocoracinae of Mayr and Amadon, 1951). Cormorants; Paleocene-Recent; Miocene-Recent in Australia; family cosmopolitan; 2-3 living, 4 extinct genera including *Graculavus* (Paleocene of North America), *Actiornis* (Eocene of England), *Pliocarbo* (Pliocene of Europe), and *Plotopterum* (Miocene of North America); one genus, *Phalacrocorax*, with a cosmopolitan distribution, only member of family in Australia; fossil record of *Phalacrocorax* extends into early to mid-Oligocene; living members of family equally diverse and endemic in New and Old Worlds; nearest relatives the specialized Anhingidae (see below); closest relatives of the

cormorants-anhingas possibly contained in the suborder Sulae (see Brodkorb, 1963; in agreement with Lanham, 1947) including the Eloptrygidae (Cretaceous-Eocene of Europe) and the Sulidae (first occurrence in the Oligocene of Europe, Miocene of North America; cosmopolitan at present).

Although *Phalacrocorax carbo* suggests by its cosmopolitan distribution that an Indomalaysian or oceanic route has been used most recently by the cormorants, direction of movement along that route is not certain. The Phalacrocoracidae are quite an old group, first recognized in the late Mesozoic/Paleogene of North America and Europe, as is *Phalacrocorax*, which is first known in the mid-Oligocene. Unfortunately, none of the practical clues used in this study favour one route over another. Phyletic relationships of the phalacrocoracids, including *Phalacrocorax*, to other birds (other than to the more specialized tropical anhingas) are not well understood, diversity and endemism are about uniform the world over, and distribution is nearly continuous along continental margins and somewhat inland over most of the world except in the far north and far south. To further complicate matters, the cormorants are not restricted to fresh-water environments, and oceanic dispersal could easily have played an important part in their dispersal with regard to Australia.

Thus, with present evidence, it appears impossible to determine the route initially used by the cormorants between Australia and the rest of the world.

Anhingidae. (Anhinginae, Mayr and Amadon, 1951). Anhingas, darters, or snake birds; ?Eocene-Recent; Miocene-Recent in Australia; 2 genera in the family (*Protoplotus*, restricted to ?Eocene of Sumatra); one genus, *Anhinga*, in Australia, has pantropical distribution; family presently with similar diversity and endemism throughout tropics of the world; nearest relatives the Phalacrocoracidae, with a cosmopolitan distribution.

The one species of anhinga (*A. melanogaster*), occurring in Australia, also extends throughout tropical Asia and tropical Africa, being distinct from the New World *A. anhinga*. A second distinct species, *A. laticeps*, was

reported from the Pleistocene of Australia (DeVis, 1905; confirmed by Miller, 1966a). Because of their present and past (see Rich, 1972, 1974) restriction to warm, wet climates, an Indomalaysian dispersal route seems most probable. At no time during the late Mesozoic or early Tertiary could the Antarctic route have had a climate that was more equable than temperate. The warm, humid tropical climate presently characteristic of some parts of northern Australia and New Guinea is a mid to late Tertiary innovation as the Australian plate drifted into the lower latitudes. The occurrence of *Protoplotus*, thought to be an early member of the Anhingidae (Lambrecht, 1931), in Paleogene sediments of Sumatra does not contradict the above conclusion.

Ardeidae. Herons and allies; early Eocene to Recent; no fossil record in Australia; family cosmopolitan; 27 living, 8 extinct genera; 9 living genera in Australia-New Guinea, none of which are endemic; one-third of genera in family with cosmopolitan distribution; greatest diversity and endemism in both New and Old Worlds, similar in both areas; nearest relatives perhaps the cosmopolitan Ciconiidae (Ligon, 1967, would disagree) (first appear in Eocene in circum-Mediterranean area; Oligocene in South America; Miocene in North America; Pliocene in Asia; Pleistocene in Australia) or the nearly cosmopolitan Threskiornithidae (first occur in Eocene in Europe; Miocene in South America; and Pleistocene in North America, Africa, and Australia).

Hérons and allies have had a long history, the family certainly being differentiated by the early Tertiary (Eocene of Europe and North America; Eocene-Oligocene of Africa). The most recent movement of this group has been across the Indomalaysian route as exemplified by the cattle egret (*Bubulcus* = *Ardeola*) that has reached Australia from Asia in the last few years (Crosby, 1972). The remaining ardeids, except for *Zonerodius* (restricted to N.G.-SW Pac.), are all wide ranging genera, most of which are decidedly more diverse outside of Australia. Diversity and endemism of the family is about equal in both the New and Old World while low in Australia, all of which suggests that the Ardeidae dispersed southward from Palearctica into Australia

and/or may have utilized an oceanic route.

Ciconiidae. Storks; Eocene-Recent; Pliocene-Recent in Australia; family cosmopolitan except for New Zealand; 11 living, 9 extinct genera only one of which (*Xenorhynchus*) is known in Australia; greatest diversity and endemism of family in Old World (Africa and Asia in particular, although three genera known in New World); nearest relatives probably Ardeidae or Threskiornithidae (see above), both of which have recent distributions that are nearly, if not, cosmopolitan and long fossil records (Eocene to Recent), or Cathartidae (Ligon, 1967).

Owing to the low diversity (one endemic genus with one species) of this family in Australia and its high diversity and endemism in neighbouring Asia (as well as the remaining Old World), entrance of the Pleistocene-Recent members of this group from Southeast Asia via the Indomalaysian route seems most probable.

Threskiornithidae. (= Plataleidae of Brodkorb, 1963). Ibises and spoonbills; Eocene-Recent; Pleistocene-Recent in Australia; 17 living, 3 extinct genera; 3 living genera in Australia; family cosmopolitan; greatest diversity and endemism in Old World with 7 genera known from New World; nearest relatives, Ciconiidae or Ardeidae, both with nearly or completely cosmopolitan distributions and long fossil histories (Eocene to Recent).

The three genera of threskiornithids represented in Australia have either cosmopolitan (*Plegadis*) or wide ranges in the Old World (*Platalea*, *Threskiornis*), and none is endemic to, or any more specifically diverse in, Australia than elsewhere. Because of such low diversity and the lack of endemic generic or suprageneric taxa in Australia at present, an Indomalaysian route for dispersal of those forms in Australia appears most probable.

Phoenicopteridae. Flamingos; late Eocene-Recent; Miocene-Pleistocene in Australia (presently extinct); nearly cosmopolitan; 3 living, 3 extinct fossil genera (*Elornis*, Eocene-Oligocene of Europe; *Tiliornis*, Oligocene of South America; *Phoeniconotius*, Miocene of Australia); no living phoenicopterids in Australia but an endemic genus (*Phoeniconotius*) and *Phoenicopterus* present during Miocene, *Phoeniconaias* in the late Pliocene, while *Phoenicop-*

terus persisted into Pleistocene; greatest diversity of family presently in South America and Africa but, when fossils considered, also in Australia; nearest relatives the Palaeodidae (Miocene-Pliocene of Holartica), Agnopteridae (Eocene-Oligocene of Europe); Telmatidae (early Eocene of South America), Scaniornithidae (early Paleocene of Europe), Torotigidae (late Cretaceous of Holartica), and more distantly the Ciconiidae and Threskiornithidae, and probably the Anseriformes (Delacour and Mayr, 1945; Hopkins, 1949) and Ciconiiformes (Mayr and Amadon, 1951; Glennys, 1955; Sibley, 1960) (both orders, Mainardi, 1962, 1963) all of which have cosmopolitan ranges but differing degrees of diversity and endemism in different parts of the world.

Although not members of the living avifauna of Australia, during the Tertiary the flamingos formed a moderately diverse group on that continent, including one endemic genus (*Phoeniconotius*). During the Pleistocene, this group became extinct in Australia (Miller, 1963b), but survived in other parts of the world in varied climatic zones from tropical to temperate. All of the Australian genera, except *Phoeniconotius*, are also represented in the Old World (*Phoeniconaias* restricted Africa), or over most of the world in tropical to temperate climates (*Phoenicopterus*). At present the family does not occur in Southeast Asia and the East Indies, being restricted further west in Eurasia and Africa in the Old World. Because the family had differentiated by the early Tertiary in both Europe and South America, and is at present equally diverse and endemic in both geographic areas, both the Indomalaysian and Antarctic dispersal routes must be considered seriously. The Indomalaysian or an oceanic route has probably been utilized most recently, accounting for the Pliocene-Pleistocene similarity at the generic level of *Phoenicopterus* (in fact Miller (1963b) suggested specific synonymy with *P. ruber* in the Pleistocene) as well as close resemblance of the Australian Pliocene *Phoeniconaias* and the same genus occurring in Africa today. Whether this was brought about strictly by a northward movement during the Miocene (or slightly before) in the case of *Phoenicopterus*

(first recorded in Europe during the early Miocene) and then or perhaps later in the case of *Phoeniconaias*, or whether movement was southwards from Eurasia, or both (different for each genus), is at present unclear. Since the extinct European Tertiary forms *Elornis* and *Tiliornis* have not been thoroughly re-examined since the late 19th century, it is uncertain how closely either is related to the Recent and other fossil genera from Europe, Africa, Australia, and the New World. At present, then, which was the initial route of dispersal between Australia and the remaining world has not been determined.

Anatidae (including Anseranatidae of Woolfenden, 1961). Ducks, geese, swans; Eocene-Recent; Miocene-Recent in Australia-New Guinea; 51 living, 23 extinct fossil genera, 13 genera in Australia-New Guinea of which *Anseranas*, *Stictonetta*, *Malacorhynchus*, *Cereopsis*, *Chenonetta*, and *Biziura* are endemic; family cosmopolitan and only slightly more diverse in Old World than elsewhere; many genera with Holarctic distribution; nearest relatives probably the Anhimidae (screamers), presently restricted to South America.

Entrance of the many different groups of Anatidae into Australia has probably been complex, possibly involving both the Antarctic and the Indomalaysian routes. Differentiation of the family occurred early, at least by the Eocene and probably earlier. The most closely related family to the anatids is the Anhimidae, present only in South America today with a Pleistocene record on that continent. In addition, many anseriforms in the Australian avifauna are primitive members of the order. The magpie geese, Anseranatidae (Woolfenden, 1961; Johnsgard, 1961, 1962; Sibley and Ahlquist, 1972) or Anseratininae of many classifications (Mayr and Amadon, 1951; Delacour, 1954), presently restricted to Australia, are anseriforms somewhat advanced over the anhimid condition but not so highly advanced as the true ducks-geese-swans (Woolfenden, 1961; Verheyen, 1953, 1955; Sibley (1960) has noted their distinctiveness from other anatids). The Cape Barren Goose (*Cereopsis*) is considered by some (Woolfenden, 1961; Johnsgard, 1961, 1962) to be the most

primitive member of the Anserinae (swan-geese subfamily). Others would disagree, however, and ally it with the sheldrakes (Tadorini) (Delacour and Mayr, 1945, 1946). *Cereopsis*, like *Anseranas*, is restricted to Australia presently, with a probable Pleistocene record of the family in New Zealand (evidencing ability of this group to cross water barriers in the past if interpretations of the New Zealand form are correct). The Dendrocygnini (tree-ducks) are considered more advanced toward the goose-swan condition than *Cereopsis* but more primitive in many characters than the Cygnini and Anserini themselves. Within the tribe of tree-ducks (Woolfenden, 1961) one very distinct genus, *Stictonetta*, is at present restricted to Australia, while *Dendrocygna* occurs pantropically, including Australia. The tribe is first recorded in Miocene sediments of South Dakota, first known in Australia in the Pleistocene. Frith (1964a-b), on the other hand, has allied this form with the swans and thus removed it from the Dendrocygnini. Since the most advanced subgroups within the Anatidae proper had differentiated by the end of the Paleogene, ancestors of more primitive members of the Anseriformes discussed above must have been present even earlier, and thus a southern dispersal of this group is a possibility. Then, too, such forms could have been relict in Australia after utilizing an Indo-malaysian route or even oceanic dispersal.

The true geese (Anserini), are presently restricted to Holarctica, apparently never reaching Australia, while the remaining tribe of the Anserinae, the swans (Cygnini) have a nearly cosmopolitan distribution. The Cygnini are first known from Oligocene sediments in Europe, from the Pleistocene elsewhere. This early occurrence in Europe coupled with the group's maximum diversity in Eurasia presently and its probable derivation from a strictly Holarctic group (Anserini) all suggest southward dispersal via the Indomalaysian route.

Members of several tribes of the Anatinae (Tadorini, Anatini, Aythiini, Oxyurini of Woolfenden (1961)) including both primitive and advanced forms are present in Australia today. This subfamily was distinct at least by the late Oligocene when it was present in

Eurasia, in Africa by the early Miocene. Primitive members of the Anserinae that probably most closely resemble the common ancestor of the Anserinae-Anatinae (Woolfenden, 1961) occur in Australia (Cercopsini) and pantropically (or nearly so: Dendrocygnini).

It is interesting that one of the more primitive tribes of the Anatinae (Woolfenden, 1961), the Oxyurini (stiff-tailed ducks), is most diverse in South America (*Heteronetta*, *Nomonyx*, *Oxyura*) while Africa and Australia each have an endemic genus (Af.-*Thalassornis*; Aust.-*Biziura*), and both share *Oxyura*, the only form present in Holarctica. Such a distribution could be explained either (1) by initial use of the Antarctic and then a northward dispersal during the Miocene into Asia and across tropical southern Asia, Europe, and North Africa into the present Ethiopian realm before restriction of the tropics in the late Tertiary, or (2) by southern dispersal into Australia from southeastern Asia in the mid-Tertiary and differentiation since that time. At present it is difficult to make a choice between the two possibilities.

Other tribes of the Anatinae, however, have apparently arrived in Australia via southward movement along the Indomalaysian route. The Tadornini (sheldrakes), have a cosmopolitan distribution, but no one genus is cosmopolitan (2 endemic to South America, 1 to Africa, 1 to Eurasia, and 2 occur over most of the Old World including Australia). Low diversity and lack of endemism in Australia, as compared to that of the Old World, indicate that the group has moved south, instead of north along the Indomalaysian route.

The Anatini are also cosmopolitan, although several genera are endemic to different parts of the world including *Malacorhynchus* (whose relationships are poorly understood, R. Schodde, pers. comm., 1974) and *Chenonetta* in Australia; *Cheniscus* is endemic to Southeast Asia-Southwest Pacific-Australia-New Guinea. Diversity and endemism in the Old and New World are nearly equal, with that in Australia being extremely low when compared with the other continents. Because of such lack of diversification in Australia in comparison to the rest of the world, the history of the Anatini

there is apparently relatively short, and again Indomalaysia seems to be the only major dispersal route utilized.

The fourth anatine tribe, the Aythyiini, is represented by only the cosmopolitan *Aythya* (by one endemic species) in Australia, a genus much more diverse elsewhere in the world. The tribe is most diverse in Asia, most endemic in the Old World, and its earliest fossil occurrence is in the early Miocene of Europe. Woolfenden (1961) believed that the Mergini (restricted both at present and in the past to Holarctica) were the nearest, more primitive relatives of the Aythyiini. All these data strongly suggest that the Aythyiini entered Australia via Indomalaysia.

In summary, the entrance of the Anatidae into Australia has probably been complex and may have involved both the Antarctic and Indomalaysian dispersal routes. The most recent exchange, which is still in progress, undoubtedly involves only the Indomalaysian route.

Accipitridae. Hawks and eagles; Eocene-Recent; Miocene-Recent in Australia; family cosmopolitan; 63 living, 24 extinct genera; 17 living genera in Australia-New Guinea of which *Lophoictinia*, *Hamirostra* (all *Milvinae*), *Erithrorhynchus* (*Accipitrinae*), and *Harpyopsis* (*Buteoninae*) are endemic; *Henicopernis* (*Perninae*) is endemic to New Guinea and the Southwest Pacific, *Haliastur* to Australia-New Guinea-Southwest Pacific-Asia, and *Butastur* to New Guinea-Asia-Africa; most closely related families *Falconidae*, *Pandionidae*, also with cosmopolitan distribution.

The Falconiformes is the most generically diverse order of non-passeriform birds, containing some 81 genera in the Peters (1931) classification and slightly less in that of Brown and Amadon (1968). Within this order, the *Accipitridae* is the most diverse family with more than 60 genera (63, Brown and Amadon, 1968; 68, Peters, 1931), equalled in intrafamilial diversity only by the parrots and pheasants. About twenty-five per cent of known accipitrid genera occur in Australia and come from several of the subfamilies within this family including three types of kites (*Perninae*, *Elaninae*, *Milvinae*), the fish eagles (*Haliaeetus*), the harriers (*Circinae*), and the hawks and eagles

(Accipitrinae). Conspicuously absent in this avifauna, however, are any of the primarily scavenging forms in the subfamily Gypaetinae, widespread in the Old World.

Most intriguing with regard to Australian biogeography are the milvine kites, thought to be some of the most primitive accipitrids (Brown and Amadon, 1968). Of a total of seven genera known, four occur in Australia, two of which (*Lophoictinia* and *Hamirostra*) are endemic to that continent. *Haliastur* occurs only in Australia - New Guinea - Southwest Pacific-Southeast Asia, while the fourth genus, *Milvus*, is found over most of the Old World in addition to Australia and the Southwest Pacific. The remaining three genera are restricted to the New World. Thus, the main diversity of this group occurs in the New World and Australia with only two genera of the seven reaching Southeast Asia and one reaching Europe. The fossils are consistent with the recent form with *Milvus* represented in the early Miocene of Europe and two presently extinct genera restricted to the New World. Certainly more advanced members within the Accipitridae had differentiated by the Eocene (Brodkorb, 1964), thus signalling an even earlier differentiation of kite ancestors. Such evidence suggests that an Antarctic route for entrance of the milvines should not be discounted. The Indomalaysian route is presently in use and could have been as early as the Miocene. Movement across this route, however, may just as well have been northwards from Australia, rather than southwards from Eurasia.

Elanus, with a nearly cosmopolitan distribution, represents the only member of Elaninae (also thought to be primitive accipitrids, Brown and Amadon, 1968) in Australia. Two other genera are in this subfamily, one genus endemic to the New World, and a second endemic to Africa. Owing to the cosmopolitan nature of *Elanus* and the lack of any more than specific differentiation of this widespread genus in Australia (*E. notatus*, *E. scriptus*), southward dispersal along the Indomalaysian route seems probable.

The third group of primitive accipitrids, the perninc kites, include seven genera, three of which occur in Australia: *Aviceda* (Af.-As.-

N.G.-Aust.-SW Pac.), *Machaerhamphus* (Af.-As.-N.G.-SW Pac.), and the Australian endemic *Henicopernis* (N.G.-SW Pac.). Three other genera are endemic to the New World and a fourth occurs in the Old World and the Southwest Pacific. As with the milvines, an Antarctic route for the subfamily should not be excluded from consideration, but neither should the Indomalaysian route. The subfamily is about equally diverse in Southeast Asia and Australia-New Guinea-Southwest Pacific, although *Aviceda* is much more diverse specifically in Southeast Asia than in Australia.

Only one genus, *Circus*, represents the Circinae in Australia. This genus has a cosmopolitan distribution, and is most diverse in the Old World, which suggests use of an Indo-malaysian dispersal route, with birds moving southward from Asia.

The Accipitrinae are represented by seven genera in Australia and New Guinea: those with (1) a nearly or entirely cosmopolitan distribution (*Accipiter*, *Spizaetus*, *Aquila*); (2) an Old World-Australian and/or New Guinean distribution (*Buteastur*, *Hieraetus*); and (3) a range restricted to Australia and/or New Guinea (*Harpyopsis*, *Erythrotriorchis*). More primitive members of the subfamily (Brown and Amadon, 1968), including the harriers and harrier hawks, are cosmopolitan. Members of the subfamily Circactinae, probably most closely related to, yet more primitive than, the accipitrines are restricted to the Old World. Since the accipitrines have very similar diversities (more than ten genera each out of a total of more than 30) in South America, Africa, and Asia but has lower diversity and little endemism in Australia, an Old World origin seems most plausible for the Australian forms. The Indomalaysian archipelago may well have been the only route utilized by accipitrines now known in Australia between that continent and the rest of the world.

Pandionidae. Ospreys; Pleistocene-Recent; only Recent in Australia; one genus (*Pandion*), which has nearly cosmopolitan distribution including Australia-New Guinea; family probably most closely related to Accipitridae (Hudson, 1948; Jollie (1953) uncertain about relationships with falconiforms; Compton

(1938) allies with Cathartidae), which likewise has cosmopolitan distribution.

The osprey (*Pandion haliaetus*) is the only member of this family and has a nearly cosmopolitan distribution, but does not occur in New Zealand. Its record extends into the Pleistocene of Europe and North America but no further. Due to its widespread nature and singular lack of diversity, it might be thought of as a relatively recent development (Ricklers and Cox, 1972; Mayr, 1963). Thus, the Indomalaysian dispersal route has most probably been involved and not the Antarctic. Whether the direction of movement was north or south can be questioned, and until the relationships of the pandionids are more clearly understood, an Australian origin cannot be ruled out.

Falconidae. Falcons/caracaras. Miocene-Recent; ?Pleistocene-Recent in Australia; family cosmopolitan; 11 living, 3 fossil genera; 1 living genus (*Falco*, nearly cosmopolitan) and possibly 1 endemic fossil genus (*Plioetus*⁹) in Australia; most closely related family, Accipitridae (also cosmopolitan in distribution) or even the Strigiformes (Sibley and Ahlquist, 1972) (Eocene-Recent, cosmopolitan).

The Falconidae, most diverse in the New World, is represented in Australia by a single living genus and possibly one endemic form, thought by most workers to be congeneric with *Falco*. *Falco* is quite diverse world-wide, with only seven species in Australia-New Guinea. Thus, it has most probably utilized the Indomalaysian route, moving southwards from Asia.

Megapodiidae. Mound builders, brush turkeys; Pleistocene-Recent; family restricted to Australia, New Guinea, and the Southwest Pacific; 7 living, 1 fossil genera; 6 living genera (2 endemic: *Leipoa* and *Alectura*) and 1 fossil (*Progura*) from Australia; family with about equal diversity and endemism in Australia-New Guinea and the Southwest Pacific; most closely related family Cracidae (curassows and chachalacas) of tropical America (Hudson *et al.*, 1966, 1969; Holman, 1964; Cracraft, 1972, 1973) although some would dispute this relationship (see refs. in Sibley and Ahlquist, 1972).

The mound-builders are galliforms endemic

⁹ Named by DeVis and needs restudy to ascertain its taxonomic position.

to Australia and the southwestern Pacific. They are thought by some workers to be primitive within this order and most closely related to the South American chachalacas (Cracidae; Cracraft, 1972, 1973; Hudson *et al.*, 1959, 1966; Chandler, 1916). Others have suggested, on the contrary, that megapodes are specialized phasianids (Clark, 1960, 1964a-b) or that cracids are more closely related to pheasants than either is to the megapodes (Sibley and Ahlquist, 1972). Providing that megapodes are actually primitive galliforms, they may be assumed to have differentiated by the Eocene (Brodkorb, 1964) because pheasants had appeared by this time. In such a case, an early southern dispersal of the mound-builders would have been a distinct possibility. On the other hand, the group could also have moved south across the Indomalaysian route and been isolated there at some time during or after the mid-Tertiary. Both hypotheses are viable but neither is more probable than the other in light of available data.

In Australia two living and one extinct (a giant Pleistocene form, *Progura* (van Tets, 1974a)) genera are endemic. Three other genera (*Megapodius*, *Telegalla*, *Aepyodius*) occur in Australia and/or New Guinea-Southwest Pacific, while *Macrocephalon* and *Eulipoa* occur only in the Southwest Pacific.

Phasianidae. (Phasianinae of Mayr and Amadon, 1951; Brodkorb, 1964). Quail, pheasants; early Oligocene-Recent; Pleistocene-Recent in Australia/New Guinea; two subfamilies—Phasianinae, restricted to the Old World and Australasia; Odontophorinae, restricted to the New World; 59 living, 8 extinct genera; 4 living genera in Australia-New Guinea, of which only *Anurophasis* is endemic (restricted to New Guinea at present but has Pleistocene record in south-eastern Australia); closely related to the Numinidae (guinea fowl, restricted to Africa presently, Pleistocene record in Europe), the Holarctic Tetraonidae (grouse, first occurs in early Miocene of North America, Pleistocene in Europe), and the North American Meleagridae (turkeys, first occur in Miocene of North America); more primitive but related to all the above are the Cracidae-Megapodiidae (see above).

The family Phasianidae is first known in the

early Oligocene of North America and Europe (different subfamilies on each continent). In Europe it was taxonomically diverse in the mid to late Cenozoic and is most diverse in Asia and islands of the Southwest Pacific today. All of the genera of phasianids occurring at present in Australia belong to the Old World subfamily, Phasianinae. One endemic genus (*Anurophasis*) has diverged from the Old World forms, but *Coturnix*, *Synoicus*, and *Excalfactoria* also occur in the Old World and Southwest Pacific. Since the Australian diversity and endemism are so low, the Asian and Southwest Pacific diversity so high, coupled with the conspecificity of three of the four Australian genera with Old World forms, dispersal southwards along the Indomalaysian route is the most probable explanation for the pheasants and quail presently known in Australia.

Turnicidae. Bustard quail; Pleistocene-Recent; 2 genera (*Turnix*, *Ortyxelos*), only *Turnix* occurs in Australia; family restricted to Old World, Southwest Pacific, and Australia/New Guinea with a fossil record in Asia and Australia; most closely related family, the Australian endemic Pedionomidae, nearest relatives to birds outside of Turnicidae-Pedionomidae uncertain (see Bock and McEvey, 1969) but most likely lie within the Gruiformes.

Pedionomidae. Collared hemipode; no fossil record; 1 genus and species present in Australia; most closely related family, the Turnicidae.

The bustard quail (Turnicidae) are generically most diverse in Africa where both genera in this family occur. *Turnix* is the only genus that reaches Australia and is itself most diverse there and in the Southwest Pacific. With this evidence alone, the most likely path of interchange between Australia and the rest of the world is Indomalaysian. But in which direction did the movement occur? The Pedionomidae, endemic to Australia, may well be the most closely related, yet perhaps more primitive group (retains hallux; Bock and McEvey, 1969), to the Turnicidae. With lack of phylogenetic analyses that point to the group of birds most closely related to the Turnicidae-Pedionomidae, it is not possible to suggest

which dispersal route(s) may have been utilized by ancestral turnicids or pedionomids. Similarity between the Turnicidae of the Old World and Australia could be accounted for by a mid-late Tertiary interchange, either northwards or southwards, along the Indomalaysian route, and the differences in diversity between Australia and Africa are not great enough to significantly favour either possibility.

Gruidae. Cranes; Eocene-Recent; Miocene-Recent in Australia; family nearly cosmopolitan except for South America and New Zealand; 4 living, 8 extinct genera; 1 genus (*Grus*) in Australia; family most diverse and with greatest endemism in Africa; nearest relatives within the Gruiformes, possibly the Aramididae (limpkins) of the New World (first recorded in Oligocene of North America (R. Emry, S. Olson, pers. comm., 1972), the Psophiidae (trumpeters, presently restricted to South America, with no fossil record), or the Eogruidae (early Oligocene of Asia (see Cracraft, 1969)), Rallidae, Heliornithidae, Eurypygidae.

The family Gruidae had differentiated by the early-mid Eocene in Holarctica and apparently was present in Australia by the mid-Miocene (Stirton, Tedford, and Woodburne, 1968). In the Recent fauna, the Gruidae are most diverse in the Old World, with only the genus *Grus* extending into the New World and Australia. *Grus* is most varied in Asia, with only two of the total nine species in the genus occurring in Australia-New Guinea (including the endemic *G. rubicuna*). Thus, based on the low endemism and diversity of the Gruidae in Australia, the high diversity in the Old World, and lack of conflicting evidence from the fossil record, southward movement over the Indomalaysian route would best explain the present record of cranes in Australia.

Rallidae. Rails; Cretaceous or Eocene-Recent¹⁰; probably mid-Miocene-Recent in Australia; family cosmopolitan; 51-52 living, 27 extinct genera; 13-14 living genera in Australia; possibly one extinct fossil genus (Stirton, *et al.*,

¹⁰ If *Telmatornis* included in the Rallidae (*vide* Brodkorb (1967)), family first known in Cretaceous; Cracraft (1969) has presented convincing arguments for placing this genus in a separate family, and thus the first record of the Rallidae is Eocene.

1968) in the mid-Miocene of central Australia; greatest diversity in Southwest Pacific region with slightly less diversity in Old World and New World; greatest number of endemic genera in South America, but nearly as many in Southwest Pacific; closest relatives of Rallidae lie within the Gruiformes, perhaps the Gruidae, Aramidae, Psophiidae, Turnicidae, Heliornithidae, Eurypygidae (see above).

At least nine genera of rallids are present in the living avifauna of Australia, of which three are endemic to Australia and/or New Guinea: *Tribonyx* (= *Gallinula* in Condon, 1973; rarely a vagrant to New Zealand), *Rallicula*, and *Megacrex*. To this list can be added *Eula-beornis* and *Gymnocrex* that are endemic to Australia-New Guinea and some islands in the Southwest Pacific. All of these genera contain only one species in Australia, except *Rallicula* with four. Three other genera range through Australia-New Guinea-Southwest Pacific-Asia (*Rallina*, *Amaurornis*, *Poliolimnas*), while the remaining forms have cosmopolitan (*Rallus*, *Porzana*, *Gallinula*, *Fulica*) or nearly cosmopolitan (*Crex*, *Porphyrio*; not found in the New World) distributions. Although genera that occur in the living avifauna of Australia have very brief (Pliocene or younger) fossil records, other members of the family are known far back into the Tertiary.

If identifications are correct, well differentiated rails were present in Europe during the Eocene and Oligocene when Australia was still attached to Antarctica. This information, coupled with the low degree of endemism and lack of speciation of endemic forms in Australia, suggests that the Indomalaysian route or a transoceanic route utilized late in the Cenozoic was probably responsible for rails presently known in Australia. The great diversity and high endemism in the Southwest Pacific is probably due to geographic isolation imposed by the insular nature of the region as well as a long history in this area, most of which has been closely associated with Asia through the Cenozoic; the diversity in South America may be due to a combination of environmental diversity as well as a long history of the group there. Perhaps the rallids were one of the groups present in South America during the

early Tertiary (no fossil record corroborates this, however) that did not successfully cross the Antarctic sweepstakes route into Australia.

Otididae. Bustards; Eocene-Recent; late Pliocene-Recent in Australia; family restricted to Old World and Australia-New Guinea; 11 genera in recent avifauna, 1 extinct fossil genus; 1 living genus (*Eupodotis* = *Ardeotis*) in Australia; most closely related family probably the extinct Gryzajidae (early Pliocene of eastern Europe); other closely related forms probably within the Gruiformes, perhaps the Cariamidae (seriemas, Pleistocene-Recent, South America).

Bustards, restricted to the Old World and Australia and particularly diverse in Africa, are first recognized in the middle Eocene of Europe. Modern genera first appear in the Miocene (*Chlamydotis*) and Pleistocene (*Otis*, *Tetrax*). Only one species of one genus (*Eupodotis australis*) is known in Australia, and its entrance seems most probably to have been via southward expansion along the Indomalaysian route perhaps as early as mid-late Pliocene.

Jacaniidae. Jacanas; Miocene-Recent; no fossil record in Australia; family pantropical; 5 living, 1 extinct genera; 1 genus (*Jacana*) living in Australia at present; greatest diversity and endemism in Old World; closest relatives probably the Phegminornithidae (early Miocene, North America); other close relatives within the Charadriiformes (possibly the Rostratulidae (Jehl, 1968), first recognized in Eocene of Europe, at present cosmopolitan except North America or the Scolopacidae (see below)).

Only one genus (and one species) of Jacaniidae is present in Australia at present. Peters (1934) recognized an endemic genus, *Irediparra* (*I. gallinacea*), for the Australian-Southwest Pacific, but Slater (1971) included this form in *Jacana*, which otherwise has a range restricted to the New World (where it has a Pleistocene to Recent record).

Restudy of the Australian form is in order, however, to determine if it really is closer to the New World or Old World forms. It appears that the Indomalaysian route was employed (with southward movement) in the initial entrance of this family into Australia because of the low diversity in Australia and the greater

diversity in the Old World. Also the present restriction of the family to the warm, humid tropics argues against the previous use of a cool, temperate high latitude dispersal route.

Haematopodidae. (Haematopodinae in Scolopacidae of Brodkorb, 1967). Oyster-catchers; Miocene-Recent; no fossil record in Australia; family cosmopolitan; 1 genus; closest relatives probably one of the following: Palaeotringinae (Cretaceous-Eocene, North America and Europe), Scolopacidae (cosmopolitan at present; first occurs in Eocene of Europe), Aphrizzidae (= Arenariidae; no fossil record, nearly cosmopolitan but not in Africa), Charadriidae (cosmopolitan, first occurs in early Oligocene of North America and Europe), Phalaropidae (Holarctic, no fossil record), Rostratulidae (see above), Burhinidae, and Recurvirostridae (see below).

The oyster-catchers are represented by two species in the only genus of the family, *Haematopus*, in the Australian avifauna. One species (*H. fuliginosus*) is endemic to Australia, while the other is nearly cosmopolitan. The group is first recognized in the early Miocene of North America. Certainly the Indomalaysian route accounts for the present cosmopolitan nature of *Haematopus*, but what the history of the family has been and where it originated are not clear. The appearance of the genus in North America in the Miocene could be owing to origin there or somewhere else, even Australia, and thus whether movement along the Indo-malaysian route of *Haematopus* has been north or south, and which route was used by the ancestor of the oyster-catchers remains to be determined.

Charadriidae. (Charadriinae of Mayr and Amadon, 1951; Brodkorb, 1967). Plovers; Oligocene-Recent; no fossil record in Australia; family cosmopolitan; 31 living, 3 extinct genera; 5 living genera in Australia, most of which are winter residents only, with one endemic genus, *Peltohyas*; greatest diversity in Old World; greatest endemism in Old World but for a single continent in South America; nearest relatives as in Haematopodidae (see above), possibly closest to the Glarecolidae (Jehl, 1968).

Four of the five genera of plovers and dotterels occurring in Australia are cosmopolitan in distribution (except *Vanellus*, which does

not occur in the New World), and all except *Vanellus* have only winter ranges in Australia. Only *Peltohyas*, with one species, is endemic to Australia. The high diversity and endemism of this group in the Old World in comparison to that of Australia suggest that the plovers most probably dispersed south along the Indomalaysian route or across oceanic barriers rather late in the Cenozoic. The Paleogene occurrence of the charadriids in Holarctica does not contradict this conclusion.

Arenariidae (often included in the Charadriidae or Scolopacidae). Turnstones; no fossil record; family cosmopolitan; 2 living genera; 1 genus (*Arenaria*) in Australia; most diverse and endemic in North America; nearest relatives possibly the Charadriidae or Phalaropodidae (Jehl, 1968) (see above).

Since the main centre of diversity (which is low) of this family is in North America and only one cosmopolitan species (*Arenaria interpres*) occurs in Australia, dispersal to Australia most likely occurred as a southward expansion of the group along the Indomalaysian route.

Scolopacidae. (Scolopacinae of Mayr and Amadon, 1951; Brodkorb, 1967). Snipe, sandpipers; Eocene-Recent; no fossil record in Australia; family cosmopolitan; 27 living, 3 extinct genera; 11 living genera in Australia; New and Old World with about equal diversity and endemism, including Australia and the Southwest Pacific; about one-third of genera in family cosmopolitan; closest relatives probably among the following: Palaeotringidae (see above), Rostratulidae (cosmopolitan except for North America, first known in Eocene of Europe); Charadriidae, Aphrizzidae, Phalaropodidae, or Haematopodidae (see above); Jehl (1968) suggests the Gallinagoninae and Calidrinae.

The Scolopacidae are first known in the Eocene of Europe. Of those genera in Australia at present, *Limosa* is first known from the Eocene¹¹ of Europe, *Numenius* from the mid-Miocene of Europe, *Bartramia* from the mid-Pliocene of North America, *Calidris* from the early Pliocene of North America, *Philomachus* from the early Pleistocene of Asia.

¹¹ All of the early Tertiary forms need re-evaluation to establish the presence of living genera during the Paleogene.

All of the genera occurring in Australia are only boreal winter visitors, most not breeding in Australia, except for *Scolopax*. Six of the eleven genera have cosmopolitan (or nearly so) distributions, two are primarily New World genera that occasionally wander to Australia, three are restricted to Australasia and the Old World.

Quite clearly the Indomalaysian route is in use at present, with many genera following this route south during the boreal winter. Some forms (*Tryngites*, *Bartramia*), however, occasionally visit Australia using neither the Antarctic nor the Indomalaysian route but cross broad oceanic barriers. What the history of the wide ranging Scolopacidae has been is uncertain, as is its place of origin. What can be deduced from the above data is that many of the genera presently in Australia have probably made use of, and are still using, the Indomalaysian route.

Recurvirostridae. (Recurvirostrinac of Mayr and Amadon, 1951). Avocets and stilts; Eocene-Recent; no fossil record in Australia; family cosmopolitan; 3 living, 2 extinct genera; all living genera occur in Australia; most diverse in Australia; one genus to Australia (*Cladorhynchus*); most closely related family probably one of the following: Phalaropodidae (see below), Jacanidae, Haematopodidae (Jehl, 1968), Charadriidae, Aphrizzidae, Scolopacidae, Rostratulidae, or Palaeotringidae (see above).

Australia is the only continent that supports all three genera (*Himantopus*, *Cladorhynchus*, *Recurvirostra*) of avocets and stilts; *Cladorhynchus* is restricted to Australia. *Himantopus* contains only one species (and that is cosmopolitan), while *Recurvirostra* contains four that are each restricted in range, one being endemic to Australia. With the fossil record of the group being as old as it is (if *Presbyornis* and *Coltonia* are accepted as members of the family (Brodkorb, 1967)), its history begins at least by the early Eocene). Combined with the above data on endemism and diversity of the group, an Antarctic route cannot be discounted for initial entrance into Australia. Similarly, a southwards dispersal along the Indomalaysian route cannot be discounted. The present cosmopolitan distribution undoubtedly is due to late

Cenozoic movement (north or south) along the Indomalaysian route or oceanic dispersal.

Phalaropodidae. (Phalaropinae of Mayr and Amadon, 1951; Brodkorb, 1967). Phalaropes; no fossil record; family cosmopolitan with breeding ranges exclusively in Holarctica, boreal winter ranges in southern hemisphere; 1 living genus; closest relatives probably one of following: Palaeotringidae, Rostratulidae, Scolopacidae, Charadriidae, Haematopodidae (see above); Jehl (1968) suggests the Tringini and Prosobonini.

Phalaropus includes three species, two of which have nearly cosmopolitan ranges and a third that breeds in the New World, spending its non-breeding season in Australia. Because the family is represented by a single genus that is neither very diverse nor endemic anywhere in the world, dispersal has probably been rather recent. In the case of *P. tricolor* that breeds in North America, neither Antarctic nor Indomalaysian routes have been utilized, but in the case of the other two species, movement along the Indomalaysian route seems the most probable explanation for present distributions. Determination of direction of movement is more difficult. Because Australia is used only as a boreal winter refuge by birds in this family, some workers have suggested that the group has originated elsewhere and expanded into Australia. Darlington (1957), however, rightly has raised objection to such interpretations (see above).

Burhinidae. Thickknees, stone curlews; Miocene-Recent; family cosmopolitan; 1-2 living genera (including *Orthorhamphus* in *Esacus*; some (Condon, 1973) consider only one genus, *Burhinus*, present) which occurs in Australia; one extinct genus (*Milnea*) from the Miocene of Europe; greatest endemism and diversity in Asia and Australia; closest relatives perhaps Jacanidae, Haematopodidae (see above; also Jehl (1968)), Glareolidae (see below), Dromadidae (coast and islands of north and west side some other family within the Charadriiformes. of Indian Ocean, no fossil record) or perhaps

The burhinids are most endemic and diverse in the Old World and Australia (*Esacus*, *Burhinus*), with only one genus (*Burhinus*) extending into the New World. Some workers

(Peters, 1934) have suggested that a third genus (*Orthorhamphus*), endemic to Australia, should be recognized. The family is not a relatively recent derivation but has a history extending at least into the early Miocene of Europe and mid-Miocene of Australia. Similarity of forms between the Old World and Australia are most probably due to dispersal across the Indomalaysian route, but what the direction of movement (north or south) has been is ambiguous. If the movement has been north, which dispersal route was employed by burhinid ancestors?

Glareolidae. Pratincoles or swallow plovers, courses; no fossil record; family restricted to the Old World-Australia-New Guinea-Southwest Pacific; 5 living genera; 2 living genera in Australia (both in the subfamily Glareolinae, the pratincoles); most diverse in Africa; one endemic genus each in Africa and Australia while remaining genera range over Old World or Old World-Australasia (excluding N.Z.); nearest relatives probably among the following: Charadriidae (Jehl, 1968), Burhinidae, Dromadidae (see above), Thinocoridae (temperate South America, no fossil record), Chionididae (southern South America and Antarctic Islands, possibly Pleistocene record in southeastern Australia, J. van Tets, pers. comm., 1972), Rostratulidae, Haematopodidae, Scolopacidae, Recurvirostridae, Phalaropidae (see above and below).

The pratincoles are represented by two genera in Australia, one endemic (*Stiltia*) and another that is widely distributed throughout the Old World (*Glareola*), though most diverse in Africa. The one species of *Glareola* (*G. maldivarum*) that occurs in Australia is only a non-breeding visitor. Closely related to the pratincoles are the Cursoriinae, considered by most workers to be the more primitive subfamily in the Glareolidae, that are entirely restricted to the Old World and particularly diverse in Africa. The family as a whole is decidedly more diverse and endemic in the Old World than in Australia, and this accompanied by the phyletic conclusions favours dispersal of the glareolids southward along the Indomalaysian route.

Laridae. (Sterninae and Larinae of Mayr and Amadon, 1951; Brodtkorb, 1967). Gulls and

terns; Oligocene¹²-Recent; probably Miocene-Recent in Australia; family cosmopolitan; 15 living, 6 extinct genera; 8 living genera in Australia; possibly 1 extinct endemic genus in Australia, described by DeVis (1905); highest diversity in New World, although not decidedly greater than in Old World; half of genera of Larinae cosmopolitan, 2 endemic to Holarctica, 1 to Galapagos; more than half of genera of Sterninae cosmopolitan, 2 endemic to South America, 2 to Australia and/or New Guinea and Southwest Pacific; nearest relatives probably Stercorariidae (skuas, oceanic birds breeding in Arctic and Antarctic, wintering on all oceans, known as fossils only from Pleistocene of North America) (Hudson *et al.*, 1969); the highly specialized Rhynchopidae (skimmers, North America, South America, Africa, and Asia, no fossil record), or perhaps birds within Brodtkorb's Scolopacidae (Palacotringinae, Rostratulinae, Scolopacinae, Aphrizinae, Charadriinae, Phalaropodinae, Haematopodinae), Recurvirostridae, Jacanidae, Burhinidae, Dromadidae, Glareolidae, Thinocoridae, Chionididae, or any other members of the Charadriiformes.

The terns are more diverse generically in Australia (as well as the rest of the world) than are the gulls. Only one genus of Larinae, *Larus*, with three species, is known in Australia, a very low diversity when compared with that of the group elsewhere in the world. In contrast, terns (Sterninae) are quite diverse, with seven of the nine genera in the subfamily represented in the Australian avifauna. Five of the seven genera are nearly, if not completely, cosmopolitan. The remaining two (*Procelsterna*, *Gygis*) are endemic to Australia and/or New Guinea and some of the Southwest Pacific islands.

Gulls are first recognized in the mid-Oligocene of Europe, and *Larus* had differentiated by the early Miocene in Europe and North America. Since the subfamily is so little differentiated and not diverse in Australia even today but is diverse outside of Australia, present members of this group most probably arrived in Australia via southward dispersal along the Indomalaysian route relatively recently. The

¹² *Halcyornis*, from the Eocene London Clay fauna, once thought to be the oldest Laridae, has recently been transferred to the Coraciiformes (Harrison and Walker, 1972).

subfamily's occurrence in the Paleogene of Europe does not argue against this conclusion.

The Sterninae are first known in Miocene sediments of Europe and occur in the mid-Miocene of Australia (Stirton, Tedford, and Woodburne, 1968). They are most diverse in Australia and the Southwest Pacific today, but most genera are cosmopolitan. Based on the subfamily's endemism, diversity, and distribution no clear decision concerning their area of origin can be made.

Thus, at present, southward movement along the Indomalaysian route most probably accounts for the presence of the known *Larinae*, but area of origin and route employed by the Sterninae is uncertain. The most recent episode of dispersal of the terns has undoubtedly been Indomalaysian or oceanic, but in what direction (north or south) and what route was utilized by the ancestors of the terns, Antarctic, Indo-malaysian, or oceanic?

Rostratulidae. (Rostratulinae of Mayr and Amadon, 1951; Brodkorb, 1967). Painted snipe; Eocene-Recent; no fossil record in Australia; family cosmopolitan except for New Zealand and North America; 2 living, 1 extinct genera; only 1 genus (*Rostratula*) in Australia; one genus endemic to South America, the other to the Old World and Australia; nearest relatives probably one of the following: Palaeotringidae, Scolopacidae, Aphrizonidae, Charadriidae, Phalaropodidae, Haematopodidae, or Jacanidae (see above).

At present a single species of painted snipe, *Rostratula bengalensis*, occurs in most of the Old World (excluding Europe) as well as in Australia and New Zealand. Although it has probably recently dispersed across the Indo-malaysian route, the direction of dispersal of this family or of its ancestors is not obvious. The only other living rostratulid is the South American *Nycticryphes*. The Recent disjunct distribution of the family suggests that previously it may have been much more widespread, or that by chance one population accidentally established itself in either South America or the Old World, while the main population remained in the homeland that spawned the founder. Because phyletic relationships have not been adequately determined for

the group, it is impossible to determine which case is more probable. In addition to the living forms, an extinct genus has been reported from the middle Eocene of Germany (which needs re-examination to ascertain its identity). If correctly identified, it has significance in signaling the differentiation of the family at an early date, for nothing is known of the group's world-wide distribution at that time. Thus, distributional history of the rostratulids is undecipherable at present; although the Indo-malaysian route has probably been used most recently, direction of that dispersal is quite uncertain.

Columbidae. Doves and pigeons; Miocene-Recent; Pleistocene-Recent in Australia; family cosmopolitan; 58 living, 3 extinct genera; 15-27 living and 1 extinct genera in Australia-New Guinea (15 living genera endemic to Aust.-N.G.-SW Pacific plus possibly endemic fossil genus described by DeVis that needs re-evaluation); most diverse and endemic in Australia and the Southwest Pacific; nearest relatives the extinct dodos and solitaires (Raphidae) of the Mascarene Islands in the Indian Ocean and possibly the sand grouse (Pteroclididae) at present occurring in much of the Old World (first recognized in the late Eocene-early Oligocene of Europe; Storer (1971) following Maclean (1967) has placed the sand grouse in the Charadriiformes; Sibley (1966), suggests relationship to both groups); relationships of the Columbiformes uncertain, perhaps showing closest affinity with the Psittacidae, more distant to the Charadriiformes or Galliformes.

At present the pigeons form a group that is extremely diverse and highly endemic in the Southwest Pacific and Australia, and only moderately so in South America, Asia, and Africa. All four subfamilies (Treroninae, Columbinae, Gourinae, Didunculinae or Ptilinopinae, Columbinae, Macropygiinae, Turturinae of Condon (1973) are represented in the Southwest Pacific and/or Australia, but only two (Treroninae, Columbinae) are represented elsewhere. The fossil record of the group is so incomplete as to be useless (except that it indicates the group had differentiated at least by the Miocene and occurred in Holarctica at that time), as is the understanding of what avian group is most closely related to the

pigeons. However, due to such great diversity and endemism at both the generic and the specific levels in Australia when compared to the pigeon faunas of the rest of the world, an origin of the group and/or a long history on that southern continent should not be discounted, nor should the possibility of an Antarctic dispersal of the group originally.

Recent movement along the Indomalaysian route accounts for generic similarity between Old World and Australian forms, but such similarity in no way demands a northern origin for the group—it demands only interchange between the two land masses. There may, in fact, have been movement both north and south along this route when individual genera are considered.

Psittacidae. Parrots; Miocene-Recent; Pleistocene-Recent in Australia; family with pan-tropical distribution at present, Tertiary representatives in Europe; 81 living, 4 extinct genera; 37 living genera in Australia of which 37 are endemic to Australia-New Guinea-Southwest Pacific; subfamilies Loriinae, Cacatuinae, Micropsittinae endemic to Australia-New Guinea-Southwest Pacific; besides 2 endemic subfamilies in New Zealand, only subfamily occurring outside of Australia (also occurs in Australia) is the Psittacinae; Condon (1973) has recently proposed the designations Cactuidae (Proboscigerinae, Calyptorhynchinae, Cacatuinae), Psittacidae (Palaeornithinae), Loriidae, Apopsittidae, Polytelidae, Platycercidae (Pezoporinae, Lathaminae, Platycercinae) for Australian parrots; greatest diversity and endemism in Australia-New Guinea-Southwest Pacific while South America has moderate diversity; nearest relatives are uncertain but may be among the Columbiformes (pigeons, sand grouse) or the Cuculiformes (turacos, cuckoos, and allies).

Although the relationships of this group to others is not sufficiently well understood, the great diversity and endemism of the group in Australia, New Guinea, and the Southwest Pacific, in comparison to the strikingly low diversity of that group in Asia and Africa (with only two and four genera respectively), does not *obviously* suggest a northern origin for the group. Instead, I believe, it strongly suggests that more serious consideration should

be given the possibility of an origin and/or early radiation on the southern continents and an Antarctic dispersal of the group or perhaps even an origin within Australia. Undoubtedly the Indomalaysian route has most recently been used in assuring similarity of the African and Asian forms with those of the Southwest Pacific and Australia.

Cuculidae. Cuckoos; Eocene-Recent; no fossil record in Australia; family cosmopolitan, 37 living and 2 extinct fossil genera; 9-10 living genera in Australia; one (*Microdynamis*) of which is endemic and 3 of which are endemic to Australia - New Guinea - Southwest Pacific (*Scythrops*, *Rhamphomantis*, *Caliechthrus*); greatest diversity of family in Southwest Pacific and Asia with twice as many endemic genera in the New World as the Old; most closely related family probably the turacos (Musophagidae) now restricted to Africa but with a Tertiary record (Eocene-Miocene) in Europe.

Cuculids are represented by 9-10 living general in Australia, only one of which is endemic. This is in marked contrast to the great diversity and high endemism of the family in the Southwest Pacific and Asia. As the nearest relatives of the cuculids are probably the musophagids (turacos), living in Africa today (and had a history in early to mid-Tertiary Europe), this group probably dispersed into Australia southwards along the Indomalaysian route. Endemism among the cuculids is highest in the New World and the group had differentiated there by the mid-Eocene; so, an Antarctic dispersal route cannot be completely ruled out but seems less likely based on known evidence owing to the group's low endemism in Australia. Because faunal interchange between North America and Europe was high as late as the early Eocene (McKenna, 1972a), cuculids might well be expected in Europe if present in North America; thus histories in both the Old and the New World have probably been long, and could account for such great diversity and endemism in these areas. Cuculids may be one group, that although present in South America at a time when the Antarctic dispersal route was open, did not have the right sweepstakes ticket!

Marchant's (1972) suggestion that the

genera *Cacomantis* and *Chrysococcyx* were derived from a common ancestor in Antarctica and then spread north into Australia and Africa respectively, does not take into account the probable pre-early Cretaceous (ca. 200 m.y. B.P.) or early Cretaceous separation of Africa and East Antarctica (see figs. 1-2) and in no way invalidates the above discussion. This timing of the Afro-Antarctic break-up is probably too far in the past to have permitted dispersal between Australia and Africa of the modern avian families, let alone closely related genera. Certainly the warm tropical vegetational belt that lined much of southern Asia and the Mediterranean area during the early and into the mid-Tertiary would have provided an excellent corridor for distribution between Australia and Africa, once Australia had moved far enough north to allow such exchange to take place (by the Miocene). Isolation imposed when such a corridor was gradually destroyed during the Neogene by increasingly cooler world-wide climate, could easily explain the *Cacomantis-Chrysococcyx* distribution seen today.

Strigidae. (Striginae of Mayr and Amadon, 1951). Owls; Eocene-Recent; no fossil record in Australia; family cosmopolitan, 28 living and 3 extinct fossil genera; 2 genera in Australia; many genera cosmopolitan with endemism nearly the same in both New and Old Worlds; group most diverse in South America, Southwest Pacific, and Asia; nearest relatives probably the Tytonidae (barn owls; Ford, 1967 *et al.*) also cosmopolitan in distribution (known as fossils from the Miocene of Europe, elsewhere only Pleistocene or later), and the Protostrigidae (Eocene, North America) and more distantly the Caprimulgiformes (see below), and perhaps the Falconiformes.

Only two genera of typical owls occur in the living Australian avifauna, while the group is quite diverse elsewhere, particularly in the Southwest Pacific, Asia, and the New World. Apparently *Ninox* (with four Australian-New Guinean species) and *Uroglaux* (1 species in New Guinea; genus restricted to New Guinea and the Southwest Pacific) or forms that gave rise to them, dispersed southwards from Asia and the Pacific Islands to Australia along

the Indomalaysian route. Otherwise, greater endemism and diversity would certainly be expected on a continent with such a small number of nocturnal avian carnivores.

Tytonidae (including the Phodilidae, which Brodkorb (1971a) isolates as a separate family; Tytoninae of Mayr and Amadon, 1951). Barn owls; Mioene-Recent; Pleistocene-Recent in Australia; family cosmopolitan; 2 living genera; 1 genus (*Tyto*) in Australia; *Tyto* cosmopolitan, but other genus, *Phodilus*, endemic to Asia and the Southwest Pacific; greatest diversity and endemism of family in Asia and the Southwest Pacific; closest relatives probably the Strigidae or the Protostrigidae (see above).

Because the greatest diversity (both generic and specific) of this group occurs in the Old World and Southwest Pacific (only *Tyto* reaches Australia and *Tyto alba* reaches the New World), southward movement through Indomalaysia seems the most likely route between Australia and the rest of the world. *Phodilus* in addition to *Tyto* occurs in the Old World; another extinct genus in the subfamily Phodilinae occurred in the Miocene of Europe.

Podargidae. Frogmouths; no fossil record; restricted to Asia-Southwest Pacific-Australia-New Guinea; 2 living genera, *Podargus* (Aust.-N.G.-SW Pac.) and *Batrachostomus* (As.-SW Pac.); nearest relatives within the Caprimulgiformes including Caprimulgidae, Aegothelidae (see below), Aegialornithidae (Eocene-Oligocene of Europe), Nyctibiidae (restricted to American tropics with a Pleistocene record in South America), or the Steatornithidae (restricted to northern South America, no fossil record).

Aegothelidae. Owlet frogmouths; Pleistocene-Recent; Mioene-Recent in Australia; at least 1 genus (*Aegotheles*); presently restricted to the Southwest Pacific-New Guinea-Australia but has Pleistocene record in New Zealand; most diverse specifically in New Guinea; nearest relatives in the Caprimulgiformes (see above, including Podargidae).

Caprimulgidae. Nightjars; Pleistocene-Recent; no fossil record in Australia; family cosmopolitan; 19 living genera; 2 genera (*Eurostopodus* and *Caprimulgus*; Condon (1973) recognized only the latter) in Australia-New Guinea; greatest diversity in the New World, particularly in South America; largest numbers of

endemics in the New World, particularly in South America; closest relatives within the Caprimulgiformes (see above).

The Caprimulgiformes, including the three previously mentioned families (Podargidae, Aegothelidae, and Caprimulgidae), are first represented in the Eocene/Oligocene of Europe by the Aegialornithidae. An aegothelid has been recovered from Miocene sediments in Australia, but the only other record of the order is in the Pleistocene from several different localities in many parts of the world. The order is composed of two families (Podargidae, Aegothelidae) endemic to Australia-New Guinea-Southwest Pacific (and Asia in the case of the Podargidae), two endemic to tropical South America (Nyctibiidae, Steatornithidae), and one with a cosmopolitan distribution (Caprimulgidae).

In the case of the cosmopolitan Caprimulgidae, entrance into Australia appears to have been via southward movement along the Indomalaysian route. Only two genera, one cosmopolitan (*Caprimulgus*) and another shared with the Old World (*Eurostopodus*), occur in Australia. The family is much more diverse with more endemic species in the Old World than in Australia, but most diverse and endemic in the New World. Such data would seem best explained by a southward dispersal along the Indomalaysian route.

Dispersal routes utilized by the other two families or their ancestors are uncertain, however. They are either entirely (Aegothelidae), or nearly (Podargidae), restricted to and most diverse in Australia-New Guinea-Southwest Pacific including New Zealand. Only the Podargidae range into Asia. What relationships they have to the Caprimulgidae and the American tropical forms (Nyctibiidae, Steatornithidae) are uncertain, and so too is the route into Australia used by both groups or their ancestors. Did they develop in Australia and/or the Southwest Pacific and spread north along the Indomalaysian route? If so, did the ancestral stock arrive originally via the Antarctic or the Indomalaysian route? Although caprimulgiforms are known only from the Eocene/Oligocene of Europe, because of generally poor records in the rest of the world at this time,

it is impossible to determine their geographic range; primitive caprimulgiforms might well have been more widespread, and if such were the case, the Antarctic route should not be dismissed.

Hemiprocnidae. Crested swifts; Pleistocene-Recent; no fossil record in Australia; 1 genus; family restricted to New Guinea-Southwest Pacific-Asia; most diverse and endemic in the Southwest Pacific; closest relatives probably the swifts (Apodidae; but Sibley, 1960, thought this due to convergence but later (Sibley and Ahlquist, 1972) noted close similarity) that have a cosmopolitan distribution (see below).

The crested swifts form a small group (one genus with three species) that is most diverse in the Southwest Pacific. It is probably most closely related to the swifts (Apodidae) that occur throughout the world. Where the group originated is uncertain, but its dispersal between Australia and the rest of the world seems intimately tied with the Indomalaysian region. At present either a northward or southward dispersal could have occurred, depending on area of origin.

Apodidae. Swifts; Eocene or Oligocene-Recent; no fossil record in Australia; family cosmopolitan; 16 living, 1 extinct genera; 3-5 living genera in Australia; diversity is low in Europe and New Zealand but about the same for all other continents; closest relatives, the Hemiprocnidae (see above), perhaps the Trochilidae (hummingbirds, with a New World, primarily South American, distribution and a Pleistocene record there; Sibley, 1960; Cohn, 1966, however, has suggested the two groups convergent); and perhaps more distantly the Caprimulgiformes (see above).

All of the genera of swifts that occur in Australia-New Guinea are monotypic or contain only a few species (*Hirundapus*, *Apus*, *Collocalia*, *Chaetura*, *Mearnsia*); only the first three are recognized by Condon (1973) and are more diverse in the Southwest Pacific and the Old World. No endemic genera occur in Australia. The fossil record indicates that the family was present in the Old World by the Eocene or Oligocene, and both *Apus* and *Collocalia* differentiated there by early Miocene. The recent distributional and diversity data

suggest that southward movement over the Indomalaysian route into Australia is most probable.

Alcedinidae. (Halcyonidae of Brodkorb, 1971a). Kingfishers; Eocene/Oligocene-Recent; no fossil record in Australia; family cosmopolitan; 14 living, 1 extinct genera; 7 living genera in the Australian-New Guinean region, one of which is endemic to New Guinea (*Clytoceyx*) and 3 of which are endemic to Australia-New Guinea-Southwest Pacific (*Melidora*, *Dacelo*, *Tanysiptera*); greatest diversity in Australia - New Guinea - Southwest Pacific region; similar endemism in Australia-Southwest Pacific and Old World regions; nearest relatives among the following: Todidae (todies of the Greater Antilles, Pleistocene record there), Motmotidae (motmots, restricted to the American tropics with a Pleistocene record there), and Meropidae (see below).

The kingfishers, first known in the Eocene or early Oligocene of Europe and later there in the Miocene, probably dispersed along the Indomalaysian route. Diversity of the group in the New World is so low (two genera in a single subfamily Cerylinae) as to suggest a rather recent arrival there, which thus argues against an Antaretic dispersal unless diversity in the past is revealed in the fossil record of the group when better known.

Meropidae. Bee eaters; Pleistocene-Recent; no fossil record in Australia; distributed over much of Old World, Southwest Pacific-Australia; 7 living genera; 1 genus (*Merops*) in Australia; greatest diversity and endemism in Africa; closest relatives among the Alcedinidae, Todidae, Motmotidae (see above), and Upupidae (hoopoes, restricted to warm parts of Old World, Pleistocene record in Old World) and Coraciidae and Bucerotidae (restricted to Old World and Australasia, with greatest diversity in Old World and Eocene-Oligocene record in Europe).

Coraciidae. Rollers; Eocene/Oligocene-Recent; no fossil record in Australia; family restricted to Old World and Australia-New Guinea-New Zealand; 5 living, 1 extinct genera; 1 genus (*Eurystomus*) in Australia; greatest diversity and endemism in Africa (including Madagascar); nearest relatives probably Bucerotidae (see below), Upupidae, Alcedinidae, Motmotidae, Todidae, or Meropidae (see above).

Bucerotidae. Hornbills; Eocene-Recent; no fossil record in Australia; family occurs in Old World-Southwest Pacific-New Guinea; 12 living, 3 extinct genera; 1 living genus (*Aceros*), reaches New Guinea, greatest endemism in Southwest Pacific; nearest relatives probably among the following: Coraciidae, Upupidae, Alcedinidae, Motmotidae, Todidae, or Meropidae (see above).

The three families mentioned above (the bee eaters, rollers, and hornbills) have in common a great diversity only in the Old World, in contrast to a single genus representation in Australia and/or New Guinea. In addition the coraciids and bucerotids have a long history (Eocene/Oligocene-Recent) in the Old World. Thus, it seems most likely that southward movement along the Indomalaysian route accounts for their presence in the Australia-New Guinea avifauna.

In summary, there are several avian families (or their ancestors) in the Australian-New Guinean fauna whose initial dispersal to or away from that continent may well have been via either an Antaretic or an Indomalaysian sweepstakes route, or in some cases via oceanic dispersal. These families include (see Table 4):

Casuariidae
 Dromornithidae
 Phalaerocoraciidae
 Phoenicopteridae
 Anatidae (in part: Anseranatidae, Cereopini, Dendrocygnini, Oxyurini)
 Accipitridae (in part: Milvinae, Perninae)
 Megapodiidae
 Pedionomidae
 Turnicidae
 Haematopodidae
 Recurvirostridae
 Burhinidae
 Laridae (in part: Sterninae)
 Rostratulidae
 Columbidae
 Psittacidae
 Podargidae
 Aegothelidae.

The remaining families not listed above have most probably moved south into Australia across the Indomalaysian route.

TABLE 4

Compilation of dispersal routes probably used between Australia and the remaining world by those non-passeriform families comprising the Australian avifauna.

INDOMALAYSIAN ROUTE (southward movement): Podicipedidae, Anhingidae, Ardeidae, Ciconiidae, Threskiornithidae, Anatidae (in part: Cygnini, Tadornini, Anatini, Aythiini), Accipitridae (in part: Elaninae, Circinae, Accipitrinae), Pandionidae (?N., ?S.), Falconidae, Phasianidae, Gruidae, Rallidae, Otidae, Jacanidae, Charadriidae, Arenariidae, Phalaropodidae, Scolopacidae, Glareolidae, Laridae (in part: Larinae), Cuculidae, Strigidae, Tytonidae, Caprimulgidae, Hemiprocnidae (?N., ?S.), Apodidae, Alcedinidae, Meropidae, Coraciidae, Bucerotidae.

ROUTE UNCERTAIN (Antarctic, Indomalaysian, and in some cases Oceanic dispersal possible): Casuariidae, Dromornithidae, Phalacrocoracidae, Pelecanidae, Phoenicopteridae, Anatidae (in part: Anseranatidae, Cereopsini, Dendrocygnini, Oxyurini), Accipitridae (in part: Milvinae, Perninae), Megapodiidae, Pedionomidae, Turnicidae, Haematopodidae, Recurvirostridae, Burhinidae, Laridae (in part: Sterninae), Rostratulidae, Columbidae, Psittacidae, Podargidae, Aegothelidae.

For many non-passeriform families (19 of the 46 families, five of these only in part) of the obligate terrestrial birds in the living, as well as the Tertiary avifauna of Australia, at least two hypotheses are available regarding route of original entrance. In many cases there is not enough information to permit a choice of one route in preference to another, and a most important point is that such a choice *cannot* be made at present, but does exist. Many of the non-passeriform families have utilized the Indomalaysian route most recently (Miocene and more recently). The mixing of the Australian and Old World avifaunas much more thoroughly than the mammalian faunas of these two areas has increased the difficulty of distinguishing Australian and Asian elements. This late Cenozoic utilization of the Indomalaysian route should not, however, be used to suggest a northern origin for all or most of the Australian non-passeriform avifauna. The striking similarity of the Australian-New Guinean fauna to that of Asia and the Southwest Pacific tells only of the latest episode in

a much longer interchange of birds between Australia and the rest of the world.

Discussion and Conclusions

Since Mayr's (1944a) classic paper suggesting that the major part of the Australian non-marine avifauna had been derived from the north, ideas concerning continental stability have radically changed. In the 1940's and as late as the 1950's, prevailing opinion among geologists, as well as biogeographers, demanded that continents had maintained their present positions with respect to one another throughout the Phanerozoic. Only with the pioneering work of Vine and Matthews (1963) on ocean floor magnetism did consensus begin shifting in favour of the currently popular theories encompassing continental drift, ocean floor spreading, and plate tectonics. Seen in the light of new and convincing evidence, Australia (including New Guinea) is no longer thought to have remained forever bounded by 45° South latitude and the equator, but instead is viewed as a truly incredible Tertiary voyager (a Noah's Ark, McKenna, 1973), moving north from a connection with the Antarctic some 50-55 m.y. ago to its present position, a journey of some 30-50 degrees of latitude. During its northward trek, profound changes in climatic regimes affecting Australia brought about the switch from a once cool, temperate vegetation covering much of the continent to the present varied and more zoned flora with the restricted cool temperate assemblages in the south, the tropical vegetation in the north (probably a Miocene innovation), and the widespread arid assemblages that characterize much of Australia today. Undoubtedly synchronous changes occurred in those faunas inhabiting the island continent, including the birds. Unfortunately, only the closing phases of Australia's avifaunal history are known, and thus such change cannot presently be documented for birds as it has been for the floras. What the late Mesozoic and early Tertiary terrestrial faunas were like is yet a mystery.

From a review (see first section of paper) of late Mesozoic and Cenozoic geologic history of Australia, other southern continents, and the ocean basins now separating them, it is

apparent that during the last 80 to 90 million years, two possible land routes (both probably sweepstakes routes in Simpson's (1940, 1953) and McKenna's (1972a-b, 1973) terminology) between Australia and the rest of the world have been available to terrestrial vertebrates: the Antarctic and the Indomalaysian. The Antarctic route was the only path not requiring dispersal across broad ocean basins that lay between Australia and the rest of the world from the late Mesozoic until the Eocene. Between that time and sometime during the Miocene, broad water barriers on the north and an increasingly hostile climate accompanied by an ever widening water barrier on the south essentially isolated Australia from the world's remaining land masses. During this period of profound isolation (that lasted perhaps as long as 20-30 million years), the avifauna, as well as the rest of the non-marine fauna, developed without much interference from the outside, only plagued, or advantaged, by the profound climatic changes that occurred as Australia drifted northward. In the mid to late Tertiary, as Australia neared its present position, the Indomalaysian route became the only major dispersal route that could be utilized by non-marine vertebrates. The Antarctic route was inactivated by the ever increasing breadth of the water barrier and the expansion of glaciation to a present continent-wide scale.

Paleoclimatological evidence suggests that the two dispersal routes had decidedly different climates throughout their entire existence. During the late Mesozoic and early Tertiary the Antarctic route was covered in many places with sizeable forests composed of *Nothofagus*, *Araucaria*, other primitive angiosperms, gymnosperms, ferns, and scouring rushes, among others, that is, a cool temperate flora much like that covering Australia during the early Tertiary. The climate was never tropical but was in no way the severe one affecting Antarctica today. The Indomalaysian route, on the other hand, has apparently supported a tropical vegetation throughout the Cenozoic, and seemingly its climate has remained much as it is today during the entire time it has served as a major route between Australia and the rest of the world.

If the modern families of birds, like the majority of modern placental mammalian families, had first developed and radiated mainly during the late Paleogene or Neogene in Holarctica, probably no choice of routes between Australia and the rest of the world used by different groups of birds would be necessary; the Indomalaysian route would be the only major stepping stone to Australia as suggested by Mayr (1944a) for the birds and by Matthew (1915) for vertebrate faunas, primarily mammals. Unlike modern placental mammalian families, however, a large number of the living avian families apparently differentiated during the Paleogene (see figure 10). Twenty of the forty-six avian families that make up the Cenozoic avifauna of Australia, had developed by the Eocene, some possibly as early as the Paleocene (Phalacrocoracidae), a few more by the Oligocene (Phasianidae, Charadriidae, Laridae), and most of the remaining families at least by the Miocene (excepting Megapodiidae, Turnicidae, Pedionomidae, Arenariidae, Phalaropodidae, Glareolidae, Podargidae, Aegothelidae, Caprimulgidae, Hemiprocnidae, and Meropidae that have only a Pleistocene or Recent record). Differentiation undoubtedly occurred sometime before the groups first appeared in the fossil record, and thus probably more than just the twenty families with lengthy records also had records extending well back into the Paleogene. So, when considering the origin of the Australian avifauna, both Indomalaysia and Antarctica should be considered as possible routes of dispersal (see figure 9).

In summary, then, because of the early differentiation of modern bird families and the availability of two dispersal routes to and from Australia, one cannot assume that most, if not all, of the Australian non-marine avifauna originated in the north. Most probable route of initial entrance of ancestors of the following families or the families themselves is impossible to determine at present: Casuariidae (including *Dromaius* and *Casuaris*), Dromornithidae, Phalacrocoracidae, Pelecanidae, Phoenicopteridae, Anatidae (in part: Anseranatidae, Cereopsini, Dendrocygnini, Oxyurini), Accipitridae (in part: Milvinae, Perninae), Megapodiidae,

Pedionomidae, Turnicidae, Haematopodidae, Scolopacidae (in part), Recurvirostridae, Burhinidae, Laridae (in part: Sterninae), Rostratulidae, Columbidae, Psittacidae, Podargidae, and Aegothelidae. Determination of the most probable route of initial dispersal for such groups will have to await further studies on avian phylogeny and a more complete fossil record from the Australian late Mesozoic and Paleogene. A point to be stressed here, however, is that such Australian birds should be recognized as *groups of uncertain origin* and should not be lumped with those forms that most probably entered Australia from *either the north or the south*. At the species level approximately 43% (as well as 19 of the 46 families present in Cenozoic of Australia) of the non-passeriform, non-marine living avifauna of Australia fall in the category 'groups of unknown origin', hardly a small segment of the Australian bird fauna! The remaining groups, on the other hand, most probably arrived via the Indomalaysian dispersal route or in a few cases as long distance travellers across the Pacific or Indian oceans (Charadriiformes in particular).

Certainly arguments using the close similarity of the Southeast Asian and Australian avifaunas as an indicator that only the Indomalaysian route has served birds dispersing between Australia and the remaining world must be questioned. Such a similarity reflects the recent, close apposition of the Palearctic and Australian continental masses and is certainly indicative of interchange, but movement of faunas could have been northward as well as southward. Such close similarity does *not* rule out the possibility of an Antarctic dispersal at some time in the past. Such southern dispersal would have occurred at a time much more distant in the past than that across Indomalaysia. The fauna that arrived in Australia via an Antarctic route would certainly have evolved during the 40 million years since their arrival there, especially as they were probably in extreme isolation for 20 million years. Owing to the climatic changes taking place as Australia drifted northward, undoubtedly there were concomitant changes, perhaps even large-scale extinctions, in the faunas that had reached Australia across

Antarctica.¹³ It was this 'changed' fauna that probably began dispersing northward across the Indomalaysian route as Asian forms spread south when the Australian 'ferryboat' (McKenna, 1973) neared its Asian 'dock'.

In conclusion, then, I would reopen consideration of the origins of the non-passeriform avifauna of Australia and call for a re-evaluation of over 40% of that fauna as data increases, particularly that regarding phylogeny of avian groups and the fossil record in Australia. Then, and only then, can probabilities be estimated that favour either an Indomalaysian or an Antarctic route for those groups in question.

Acknowledgements

Many people are to be thanked for thought-provoking discussions that enhanced this paper, in particular W. Bock, M. McKenna, and R. Tedford, as well as D. Bohaska, J. Cracraft, I. Dalziel, T. Darragh, D. Dorwood, D. Elliot, E. Gaffney, W. Hamilton, G. and J. Hope, E. Kemp, A. McEvey, B. Nielsen, S. Olson, M. Plane, P. Raven, T. Rich, B. Schaeffer, R. Schodde, J. van Tets, and J. Warren. R. Gooris rendered figures 7, 8 and 10 and H. Galiano figures 9A-B. Financial support for this study was provided by an American Association of University Women Fellowship, an N.D.E.A. Title IV Fellowship, and an Australian-American Educational Foundation Grant for 1973-1974. Research facilities for the writing of this paper were kindly provided by The American Museum of Natural History, New York, The National Museum of Victoria, Melbourne, and The Bureau of Mineral Resources, Canberra.

¹³ Such severe climatic changes taking place in the Australian Tertiary may serve as yet another factor, in addition to those mentioned by Keast (1972), that accounts for the differences in the diversity and endemism characteristic of the South American, African, and Australian avifaunas. Both the South American and the African land masses have remained at nearly the same latitudinal positions throughout the Tertiary with concomitant stability of climatic regimes affecting those continents. Australia, on the other hand, has quite noticeably moved some 30-50° of latitude northwards. Perhaps such changes, and possibly accompanying extinctions, account in part for the lower endemism in Australia when compared to that characterizing South America as well as such factors as smaller land area, less topographic diversity, decidedly smaller areas of tropical environments, etc. suggested by Keast.

Bibliography

- ADIE, R. J., 1953. The Rocks of Graham Land. Ph.D. thesis, Univ. of Cambridge, 259 pp. [unpublished].
- ALLAN, T. D., 1969. A review of marine geomagnetism. *Earth-Sci. Rev.*, 5, p. 217-254.
- ANDERSON, M. A. and H. O. FLETCHER, 1934. The Cuddie Springs bone bed. *Aust. Mus. Mag.*, 5, p. 152-158.
- ANDREWS, H. N., 1961. *Studies in Paleobotany*. John Wiley and Sons, Inc., 487 pp.
- ANONYMOUS, 1959. A new fossil penguin from Australia. *Nature*, 184, p. 1612.
- AUDLEY-CHARLES, M. G., D. J. CARTER, and J. S. MILSON, 1972. Tectonic Development of Eastern Indonesia in Relation to Gondwanaland Dispersal. *Nature, Phy. Sci.*, 239 (90), p. 35-39.
- AXELROD, D. I., 1960. The Evolution of flowering plants. In: *Ed. Sol Tax, The Evolution of Life*. Univ. Chicago Press, Chicago, p. 227-306.
- AXELROD, D. I. and H. P. BAILEY, 1968. Cretaceous dinosaur extinction. *Evolution*, 22, p. 595-611.
- BARBOUR, G. B., 1929. The geology of the Kalgan area. *Mem. Geol. Surv. China, Ser. A*, (56), 148 pp.
- BECK, M. E., 1972. Palaeomagnetism and magnetic polarity zones in the Jurassic Dufek Intrusion, Pensacola Mountains, Antarctica. *Geophys. Jour. Roy. Astro. Soc.*, 28, p. 49-63.
- BEN-AVRAHAM, Z. and S. UYEDA, 1973. The evolution of the China Basin and the Mesozoic paleogeography of Borneo. *Earth Planet. Sci. Lett.*, 18, p. 365-376.
- BERGGREN, W. A., 1969. Cenozoic chronostratigraphy, planktonic foraminiferal zonation and the radiometric time scale. *Nature*, 224, p. 1072-1075.
- , 1971. Neogene stratigraphy, planktonic foraminiferal zonation and the radiometric time scale. *Hungar. Geol. Soc. Bull.*, 101, p. 162-169.
- BIBBY, J. S., 1966. The stratigraphy of part of northeast Graham Land and the James Ross Island Group. *Brit. Antarc. Surv., Sci. Rep.* (53), London, 37 pp.
- BOCK, W. J., 1956. A generic review of the family Ardeidae. *Am. Mus. Novitates*, 1779, 49 pp.
- , 1963. The cranial evidence for Ratite affinities. *Proc. XIIIth Internat. Ornith. Congress*, 1, p. 39-54.
- BOCK, W. J. and A. MCEVEY, 1969. Osteology of *Pedionomus torquatus* (Aves: Pedionomidae) and its allies. *Proc. Roy. Soc. Vict., N. S.*, 82(2), p. 187-232.
- BRODKORB, P., 1963. Catalogue of fossil birds. *Bull. Florida State Mus., Biol. Sci.*, 7(4), p. 179-293.
- , 1964. Catalogue of fossil birds: Pt. 2 (Anseriformes through Galliformes). *Bull. Florida State Mus. Biol. Sci.*, 8(3), p. 195-335.
- , 1967. Catalogue of fossil birds: Pt. 3 (Ralliformes, Ichthyornithiformes, Charadriiformes). *Bull. Florida State Mus., Biol. Sci.*, 11(3), p. 99-220.
- , 1971a. Catalogue of fossil birds: Pt. 4 (Columbiformes through Piciformes). *Bull. Florida State Mus., Biol. Sci.*, 15(4), p. 163-266.
- , 1971b. Origin and evolution of birds. In: *Farner, D. S. and J. R. King (ed.), Avian Biology*, 1, Academic Press, New York, p. 19-55.
- BROWN, D. A., K. S. W. CAMPBELL, and D. A. W. CROOK, 1968. The geographical evolution of Australia and New Zealand. Pergamon Press, N.Y., 409 pp.
- BROWN, L. and D. AMADON, 1968. *Eagles, Hawks, and Falcons of the World*. 2 vols. McGraw-Hill Book Co., New York, 944 pp.
- BURBIDGE, N. T., 1960. The Phytogeography of the Australian Region. *Aust. Jour. Bot.*, 8, p. 75-212.
- BURKE, K., T. F. J. DESSAUVAGIE, and A. J. WHITEMAN, 1971. Opening of the Gulf of Guinea and geological history of the Benue Depression and Niger Delta. *Nature, Phys. Sci.*, 233, p. 51-55.
- BUTLER, W. H., 1969. Remains of *Sarcophilus* the 'Tasmanian' Devil (Marsupialia, Dasyuridae) from coastal dunes South of Scott River, Western Australia. *West. Aust. Nat.*, 11(4), p. 87-88.
- CARTER, R. M. and C. A. LANDIS, 1972. Correlative Oligocene unconformities in southern Australia. *Nature, Phys. Sci.*, 237, p. 12-13.
- CHANDLER, A. C., 1916. A study of structure of feathers, with reference to their taxonomic significance. *Univ. Calif. Publ. Zool.*, 13, p. 243-446.
- CHANEY, R., T. TANAI, N. SUZUKI, *et al.*, 1963. Tertiary floras of Japan. The Collaborating Association to Commemorate the 80th Anniversary of the Geological Survey of Japan, Tokyo, 262 pp.
- CHAPMAN, A. L. S., 1910. New or little known Victorian fossils in the National Museum. *Proc. Roy. Soc. Vict., N.S.*, 23 (1), p. 21-26.
- CHRISTOFFEL, D. A. and D. I. ROSS, 1970. A fracture zone in the southwest Pacific basin south of New Zealand and its implications for sea floor spreading. *Earth Planet. Sci. Lett.*, 8, p. 125-130.
- CLARK, G. A., Jr, 1960. Notes on the embryology and evolution of the megapodes (Aves: Galliformes). *Postilla*, 45, p. 1-7.
- , 1964a. Ontogeny and evolution in the megapodes (Aves: Galliformes). *Postilla*, 78, p. 1-37.
- , 1964b. Life histories and the evolution of megapodes. *Living Bird*, 3, p. 149-167.
- COHN, J. W., 1966. Convergence between Apodi and Trochili. XIV Internat. Ornith. Cong., Abs., Oxford, p. 44-45.
- COLANI, M., 1917. Essai sur les Flores tertiaires du Tonkin. *Bull. du Service geol. de l'Indochine*, 4, 142 pp.
- , 1920. Etude sur les Flores tertiaires de quelques gisements de lignite de l'Indochine et du Yunnan. *Bull. du Service geol. de l'Indochine*, 8, 524 pp.
- COMPTON, L. V., 1938. The pterylosis of the Falconiformes with special attention to the taxonomic position of the osprey. *Univ. California Publ. Zool.*, 42, p. 173-212.
- CONDON, H. T., 1973 (in press). A synoptic list of the birds of Australia. Part I (Non-passerines). Species Nos. 1-393, Introductions and fossils.
- CRACRAFT, J., 1969. Systematics and evolution of the Gruiformes (Class Aves). 1. The Eocene family Geranoididae and the early history of the Gruiformes. *Am. Mus. Novitates*, 2388, 41 pp.
- , 1972. Continental drift and Australian avian biogeography. *Emu*, 72(4), p. 171-174.
- , 1973. Continental drift, paleoclimatology, and the biogeography of birds. *Jour. Zoology, London*, 169, p. 455-545.

- CRACRAFT, J. and P. V. RICH, 1972. Systematics and evolution of the Old World Cathartidae. *Condor*, 74(3), p. 272-283.
- CRANWELL, L. M., 1959. Fossil pollen from Seymour Island, Antarctica. *Nature*, 184(4701), p. 1782-1785.
- CRANWELL, L. M., H. J. HARRINGTON, and I. G. SPEDEN, 1960. Lower Tertiary microfossils from McMurdo Sound, Antarctica. *Nature*, 186, p. 700-702.
- CRAWFORD, A. R., 1971. Gondwanaland and the growth of India. *Jour. Geol. Soc. India*, 12(3), p. 205-221.
- CREER, K. M., 1970. A review of paleomagnetism. *Earth-Sci. Rev.*, 6(6), p. 369-466.
- CREER, K. M., B. J. J. EMBLETON, and D. A. VALENCIO, 1969. Comparison between the upper Palaeozoic and Mesozoic palaeomagnetic poles for South America, Africa and Australia. *Earth Planet. Sci. Let.*, 7, p. 288-292.
- CRIBB, H. G. S., N. R. MCTAGGERT, and H. R. E. STAINES, 1960. Sediments east of the Great Divide. In: Hill, D. and A. K. Denmead (ed.), *The Geology of Queensland*, *Jour. Geol. Soc. Aust.*, 7, p. 345-355.
- CROSBY, G. T., 1972. Spread of the cattle egret in the western hemisphere. *Bird-banding*, 43(3), p. 205-212.
- DALZIEL, I. W. D. and D. H. ELLIOT, 1971. Evolution of the Scotia Arc. *Nature*, 233, p. 246-252.
- DALZIEL, I. W. D. and D. H. ELLIOT, 1973. The Scotia Arc and Antarctic margin. In: A. E. M. Nairn and F. G. Stehli (ed.), *The ocean basins and margins*, 1, p. 171-246.
- DARLINGTON, P. J., 1957. *Zoogeography: the geographical distribution of animals*. John Wiley and Sons, London, New York, 675 pp.
- DARRAGH, T. A., 1973. (Abstr.) Upper Eocene Mollusca from North Walpole, W.A. Their stratigraphic and paleogeographic significance. 45th A.N.Z.A.S. Congress. Sect. 3, p. 9.14-9.15.
- , 1974 (ms.). Tertiary molluscan assemblages from southeastern Australia and their biostratigraphic significance. Ph.D. thesis in progress, Univ. Melbourne.
- DAVID, T. W. E., 1950. *The Geology of the Commonwealth of Australia 1*. Edward Arnold and Co., London, 747 pp.
- DAVIES, H. L. and I. E. SMITH, 1971. Geology of Eastern Papua. *Bull. Geol. Soc. Amer.*, 82, p. 3299-3312.
- DELACOUR, J., 1954. *The Waterfowl of the World. 1*. Country Life Limited, London, 284 pp.
- DELACOUR, J. and E. MAYR, 1945. The family Anatidae. *Wilson Bull.*, 57, p. 3-55.
- , 1946. Supplementary notes on the family Anatidae. *Wilson Bull.*, 58, p. 104-110.
- DEVIS, C. W., 1888a. A glimpse of the post-Tertiary avifauna of Queensland. *Linn. Soc. N.S.W.*, 2nd Ser., 3(3-4), p. 1277-1292.
- , 1888b. Australian ancestry of the Crowned Pigeon of New Guinea. *Proc. Roy. Soc. Queensland*, 5, p. 127-131.
- , 1889a. Additions to the list of fossil birds. *Proc. Roy. Soc. Queensland*, 6, p. 55-58.
- , 1889b. On a bone of an extinct eagle. *Proc. Roy. Soc. Queensland*, 6, p. 161-162.
- , 1891a. Note on an extinct eagle. *Proc. Linn. Soc. New South Wales*, 6(2), p. 123-125.
- , 1891b. On the trail of an extinct bird. *Proc. Linn. Soc. New South Wales*, 6(2), p. 117-122.
- , 1891c. Residue of the extinct birds of Queensland as yet detected. *Proc. Linn. Soc. New South Wales*, 6(2), p. 437-456.
- , 1905. A contribution to the knowledge of the extinct avifauna of Australia. *Ann. Queensland Mus.*, 6, p. 1-25.
- DICKSON, G. O., W. C. PITMAN, III, and J. R. HEIRTZLER, 1968. Magnetic anomalies in the South Atlantic and ocean floor spreading. *Jour. Geophys. Res.*, 73(6), p. 2087-2100.
- DIETZ, R. S. and J. C. HOLDEN, 1970. Reconstruction of Pangaea: Breakup and dispersion of continents, Permian to Present. *Jour. Geophys. Res.*, 75(26), p. 4939-4956.
- DIETZ, R. S. and W. P. SPROLL, 1970. Fit between Africa and Antarctica: A continental drift reconstruction. *Science*, 167(3925), p. 1612-1614.
- DINGLE, R. V., 1973. Mesozoic palaeogeography of the southern Cape, South Africa. *Palaeogeog., Palaeoclim., Palaeoecol.*, 13(3), p. 203-214.
- DINGLE, R. V. and H. C. KLINGER, 1971. Significance of upper Jurassic sediments in the Knysna Outlier (Cape Province) for timing the breakup of Gondwanaland. *Nature Phys. Sci.*, 232, p. 37-38.
- DORMAN, F. H., 1966. Australian Tertiary palaeotemperatures. *Jour. Geol.*, 74(1), p. 49-61.
- DORMAN, F. H. and E. D. GILL, 1959a. Oxygen isotope paleotemperature determinations of Australian Cainozoic fossils. *Science*, 130, p. 1576.
- , 1959b. Oxygen isotope paleotemperature measurements on Australian fossils. *Proc. Roy. Soc. Vic.*, 71, p. 73-98.
- DUN, W. S., 1894-95. Notes on the occurrence of monotreme remains in the Pliocene of New South Wales. *Rec. Geol. Surv. New South Wales*, 4, p. 118-126.
- EDWARDS, T. N., 1923. On some Tertiary plants from south-east Burma. *Geol. Mag.*, 60, p. 159-165.
- ELLIOT, D. H., 1972 (in press). Aspects of Antarctic geology and drift reconstructions. *Contrib. Institut. Polar Studies*, Ohio State Univ., Columbus.
- ENDO, S., 1964. Some older Tertiary plants from northern Thailand. In: Kobayashi, T. and R. Toriyama, *Geol. and Palaeon. Southeast Asia*, p. 113-119.
- ENDO, S. and I. FUJIYAMA, 1966. Some late Mesozoic and late Tertiary plants and a fossil insect from Thailand. In: Kobayashi, T. and R. Toriyama, *Geol. and Palaeon. Southeast Asia*, 2, p. 301-308.
- ETHERIDGE, R., 1889. On further evidence of a large extinct struthious bird (*Dromornis* Owen) from the post Tertiary deposits of Queensland. *Rec. Geol. Surv. New South Wales*, p. 126-136.
- , 1894. Vertebrate remains from the Warburton or Diamantina River. Official contributions to the palaeontology of South Australia. No. 6. *Ann. Rep. Govt. Geol. S. Australia*, 1894, p. 19-22.
- FINLAYSON, H. H., 1938. On the occurrence of a fossil penguin in Miocene beds in South Australia. *Trans. Roy. Soc. S. Aust.*, 62(1), p. 14-17.

- FLEMING, C. A., 1962. New Zealand biogeography. A paleontologist's approach. *Tuatara*, 10(2), p. 53-108.
- FLORES, G., 1970. Suggested origin of the Mozambique Channel. *Trans. Geol. Soc. S. Africa*, 73, p. 1-16.
- FORD, N. L., 1967. A systematic study of the owls based on comparative osteology. Ph.D. disser., Univ. Michigan, Ann Arbor. Abs. In: *Diss. Abs., Sec. B.*, 28(12), p. 5240.
- FRANCHETEAU, J. and J. G. SCLATER, 1969. Paleomagnetism of the southern continents and plate tectonics. *Earth Planet. Sci. Lett.*, 6, p. 93-106.
- FRITH, H. J., 1964a. Taxonomic relationships of *Stictonetta naevosa* (Gould). *Nature*, 202(4939), p. 1352-1353.
- , 1964b. The downy young of the Freckled Duck *Stictonetta naevosa*. *Emu*, 64(1), p. 42-47.
- FUNNELL, B. M. and A. G. SMITH, 1968. Opening of the Atlantic Ocean. *Nature*, 219, p. 1328-1333.
- GEITZENAUER, K. R., S. V. MARGOLIS, and D. E. EDWARDS, 1968. Evidence consistent with Eocene glaciation in a South Pacific deep sea sedimentary core. *Earth Planet. Sci. Lett.*, 4, p. 173-177.
- GENTILLI J., 1949. Foundations of Australian bird geography. *Emu*, 49, p. 85-129.
- GERMERAAD, J. H., C. A. HOPPING, and J. MULLER, 1968. Palynology of Tertiary sediments from tropical areas. *Rev. Palaeobot. and Palynology*, 6(3/4), p. 189-348.
- GILL, E. D., 1959a. Penguins—past and present. *Victorian Nat.*, 75, p. 178-179.
- , 1959b. Provenance of fossil penguin from Western Victoria. *Proc. Roy. Soc. Vict.*, 71(2), p. 121-123.
- , 1961. The climates of Gondwanaland in Kainozoic times. In: Nairn, A. E. M., *Descriptive paleoclimatology*, Interscience, New York, p. 332-353.
- , 1914. The Mammoth Cave (con'd.). *Rec. Western Australian Mus. Art Gall.*, 1(3), p. 244-251.
- , 1948. The cave fossils of the southwest. *West. Australia Nat.*, 1(5), p. 101-104.
- GLENNY, F. H., 1955. Modifications of pattern in the aortic arch system of birds and their phylogenetic significance. *Proc. U.S. Natl. Mus.*, 104, p. 525-621.
- , 1965. Main cervical and thoracic arteries of some flightless birds. *Ann. Zool.*, 5(1), p. 1-8.
- GOLENPAUL, D., 1960. *Information Please Almanac*. McGraw-Hill, New York, 912 pp.
- GÖPPERT, H. R., 1854. *Die Tertiärflora auf der Insel Java*. C. W. Mieling, S'Gravenhage, 169 pp.
- GRABAU, A. W., 1926. A summary of the Cenozoic and Psychozoic deposits with special reference to Asia. *Bull. Geol. Soc. China*, 6(2), p. 151-194.
- GRAM, R., 1969. Grain surface features in Eltanin cores and Antarctic glaciation. *Antarct. Jour.*, 6(5), p. 174-175.
- GREGORY, J. W., 1906. *The dead heart of Australia*. London, xvi + 384 pp.
- GREEN, A. G., 1972. Seafloor spreading in the Mozambique Channel. *Nature Phys. Sci.*, 236(63), p. 19-21.
- GRIFFITHS, J. R., 1971. Reconstruction of the southwest Pacific margin of Gondwanaland. *Nature*, 234, p. 203-207.
- GRIFFITHS, J. R. and R. VARNE, 1972. Evolution of the Tasman Sea, Macquarie Ridge and Alpine Fault. *Nature Phys. Sci.*, 235(57), p. 83-86.
- GRIGG, G. C., 1964. Studies on the Queensland lungfish, *Neoceratodus forsteri* (Kreffit). II. Thermal acclimation. *Australian Jour. Zool.*, 13, p. 407-411.
- HAILE, N. S. and M. W. McELHINNY, 1972. The potential value of paleomagnetic studies in restraining romantic speculation about the geological history of Southeast Asia. *Geol. Soc. Malaysia, Reg. Conf. Geol. S.E. Asia, Abstr., Annex to Newsletter*, 34, p. 16.
- HAMILTON, W., 1964. Antarctica and Pacific tectonics. *Rcpt. 22nd session, Inter. Geol. Cong. India, Pt. IV, Proc. Sect. 4*, p. 443-459.
- , 1967. Tectonics of Antarctica. *Tectonophys.*, 4, p. 555-568.
- , 1972a. Plate tectonics of Southeast Asia and Indonesia. *Geol. Soc. Malaysia, Reg. Conf. Geol. S.E. Asia, Abstr., Annex to Newsletter*, 34, p. 19.
- , 1972b. Tectonics of the Indonesian Region. U.S. Dept. Interior, Project Report, Indonesian Investigations (IR) IND-20, 13 pp.
- HARRINGTON, H. J., K. L. BURNS, and B. R. THOMPSON, 1973. Gambier-Beaconsfield and Gambier-Sorell fracture zones and the movement of plates in the Australia-Antarctica-New Zealand region. *Nature Phys. Sci.*, 245(146), p. 109-112.
- HARRISON, C. J. O. and C. A. WALKER, 1972. The affinities of *Halcyornis* from the lower Eocene. *Bull. Brit. Mus. (Nat. Hist.)*, Geol., 21(4), London, p. 153-169.
- HAYES, D. E., *et al.*, 1973. Leg 28 Deep Sea drilling in the southern ocean. *Geotimes*, June, pp. 19-24.
- HEIRTZLER, J. R. and R. H. BURROUGHS, 1971. Madagascar's paleoposition: new data from the Mozambique Channel. *Science*, 174, p. 488-490.
- HEIRTZLER, J. R., *et al.*, 1968. Marine magnetic anomalies, geomagnetic field reversals and motions of the ocean floor and continents. *Jour. Geophys. Res.*, 73(6), p. 2119-2136.
- , 1973. Age of the floor of the eastern Indian Ocean. *Science*, 180(4089), p. 952-954.
- HENNIG, W., 1966. *Phylogenetic systematics*. Univ. Illinois Press, Urbana, 263 pp.
- HILL, D. and A. K. DENMEAD (ed.), 1960. *The geology of Queensland*. *Jour. Geol. Soc. Australia*, 7, 474 pp.
- HOLDEN, R., 1916. A fossil wood from Burma. *Rec. Geol. Surv. India*, 47, p. 267-272.
- HOLMAN, J. A., 1964. Osteology of gallinaceous birds. *Quart. Jour. Florida Acad. Sci.*, 27(3), p. 230-252.
- HOPKINS, G. H. E., 1949. Some factors which have modified the phylogenetic relationship between parasite and host in the Mallophaga. *Proc. Linn. Soc. Lond.*, 161, p. 37-39.
- HOUSTON, B. R., 1967. *Geology of the city of Brisbane*. *Geol. Surv. Queensland, Pub. 324*, 86 pp.
- HUDSON, G. E., 1948. Studies on the muscles of the pelvic appendage in birds. II: the heterogeneous order Falconiformes. *Am. Mid. Nat.*, 39, p. 102-127.
- HUDSON, G. E., R. A. PARKER, J. V. BERGE, and P. J. LANZILLOTTI, 1966. A numerical analysis of the modifications of appendicular muscles in

- various genera of gallinaceous birds. *Am. Mid. Nat.*, 76(1), p. 1-73.
- HUDSON, G. E., K. M. HOFF, J. V. BERGE, and E. C. TRIVETTE, 1969. A numerical study of the wing and leg muscles of the Lari and Alcae. *Ibis*, 111, p. 459-524.
- JARDINE, N. and D. MCKENZIE, 1972. Continental drift and the dispersal and evolution of organisms. *Nature*, 235(5332), p. 20-24.
- JEHL, J. R., Jr., 1968. Relationships in the Charadrii (shorebirds): a taxonomic study based on colour patterns of the downy young. *Mcm. San Diego Soc. Nat. Hist.*, 3, p. 1-54.
- , 1971. The colour patterns of downy young ratites and tinamous. *Trans. San Diego Soc. Nat. Hist.*, 16, p. 291-301.
- JOHNSGARD, P. A., 1961. The taxonomy of the Anatidae—a behavioural analysis. *Ibis*, 103a, p. 71-85.
- , 1962. Evolutionary trends in the behaviour and morphology of the Anatidae. *Wildfowl Trust*, 13th An. Rep., p. 130-148.
- JOLLIE, M., 1953. Are the Falconiformes a monophyletic group? *Ibis*, 95, p. 369-371.
- JONES, C. R., D. J. GOBBETT, and T. KOBAYASHI, 1966. Summary of fossil record in Malaya and Singapore, 1900-1965. In: Kobayashi, T. and R. Toriyama, *Geol. and Palaeon. Southeast Asia*, 2, p. 309-359.
- JONES, J. G., 1971. Australia's Cenozoic drift. *Nature*, 230, p. 237-239.
- , 1972. Significance of upper Jurassic sediments in the Knysna Outlier (Cape Province). *Nature Phys. Sci.*, 235(55), p. 59-60.
- JONES, L. J., 1940. The Gulgong goldfield. *Dept. Mines, Geol. Surv. New South Wales. Min. Res.*, 38, p. 1-134.
- KATILI, J. A., 1971. A review of the geotectonic theories and tectonic maps of Indonesia. *Earth Sci. Rev.*, 7, p. 143-163.
- KATZ, H. R., 1973. Contrasts in tectonic evolution of orogenic belts in the south-east Pacific. *Jour. Roy. Soc. New Zealand*, 3(3), p. 333-362.
- , 1972. Faunal elements and evolutionary patterns: some comparisons between the continental avifaunas of Africa, South America, and Australia. *Proc. XV Intern. Ornith. Cong.*, p. 594-622.
- KEMP, E. M., 1972a. Recycled palynomorphs in continental shelf sediments from Antarctica. *Antarct. Jour.*, 7(5), p. 190-191.
- , 1972b. Reworked palynomorphs from West ice shelf area, east Antarctica, and their possible geological and palaeoclimatological significance. *Marine Geol.*, 13(3), p. 145-157.
- KENNETT, J. P., *et al.*, 1972. Australian-Antarctic continental drift, palaeocirculation changes and Oligocene deep-sea erosion. *Nature, Phys. Sci.*, 239, p. 51-55.
- KENNETT, J. P. and C. A. BRUNNER, 1973. Antarctic late Cenozoic glaciation: evidence for initiation of ice-rafting and inferred increased bottom-water activity. *Geol. Soc. Amer., Bull.*, 84, p. 2043-2052.
- KENT, P. E., 1972. Mesozoic history of the east coast of Africa. *Nature*, 238, p. 147-148.
- KON'NO, E., 1966. Some younger Mesozoic plants from Malaya. In: Kobayashi, T. and R. Toriyama, *Geol. and Palaeon. Southeast Asia*, 3, p. 309-359.
- KRÄUSEL, R., 1922 (not seen). Fossile Hölzer aus dem Tertiär von Süd-Sumatra. *Verhand. Geol.—Mijnb. Genootschap voor Nederlanden Koloniën (Geol. Scr.)*, 5.
- , 1925 (not seen). Der Stand unserer Kenntnisse von der Tertiär flora Niederländisch—Indiens. *Verhand. Geol.—Mijnb. Genootschap Nederland. Koloniën (Geol. Ser.)*, 8.
- , 1926. Über einige fossile Hölzer aus Java. *Leidsche Geol. Meded.*, 2, p. 1-6.
- KUMMEL, B., 1961. *History of the Earth*. Freeman and Co., San Francisco, London, 610 pp.
- LAMBRECHT, K., 1931. *Protoplopus beauforti* n.g., n. sp., ein schlangenhalsvögel aus dem Tertiär von W. Sumatra. *Wet. Meded. Dienst. Mijnb. E. Indies*, 17, p. 15-24.
- LANHAM, U. N., 1947. Notes on the phylogeny of the Pelecaniformes. *Auk*, 64, p. 65-70.
- LARSON, E. E. and L. LAFOUNTAIN, 1970. Timing of the breakup of the continents around the Atlantic as determined by paleomagnetism. *Earth Planet. Sci. Lett.*, 8, p. 341-351.
- LEMASURIER, W. E., 1970. Volcanic evidence for early Tertiary glaciation in Marie Byrd Land. *Antact. Jour.*, 5(5), p. 154-155.
- LEYDEN, R., G. BRYAN, and M. EWING, 1972. Geophysical reconnaissance on African Shelf: 2. Margin sediments from Gulf of Guinea to Walvis Ridge. *Am. Assoc. Pet. Geol. Bull.*, 56(4), p. 682-693.
- LEPICHON, X. and J. R. HEIRTZLER, 1968. Magnetic anomalies in the Indian Ocean and sea-floor spreading. *Jour. Geophys. Res.*, 73, p. 2101-2177.
- LIGON, J. D., 1967. Relationships of the cathartid vultures. *Occas. Pap., Univ. Michigan Mus. Zool.*, 651, p. 1-26.
- LONGMAN, H. A., 1945. Fossil vertebrates from Gore quarries. *Mem. Queensland Mus.*, 12, p. 164.
- MACLEAN, G. L., 1967. Die systematische Stellung der Flughühner (Pteroclididae). *Jour. Ornith.*, 108, p. 203-217.
- MAINARDI, D., 1962. Immunological data on the phylogenetic relationships and taxonomic position of flamingos (Phoenicopteridae). *Ibis*, 104, p. 426-428.
- , 1963. Immunological distances and phyletic relationships in birds. *Proc. XIII Internat. Cong.*, p. 103-114.
- MANDRA, Y. T., 1969. Silicoflagellates: A new tool for the study of Antarctic Tertiary climates. *Antarct. Jour.*, 4(5), p. 172-174.
- , 1971. Upper Eocene silicoflagellates from New Zealand. *Antarct. Jour.*, 6(5), p. 177-178.
- MANDRA, Y. T. and H. MANDRA, 1970. Antarctic Tertiary marine climate based on silicoflagellates. *Antarct. Jour.*, 5(5), p. 178-180.
- MANTEN, A. A. (ed.), 1969. *Spec. Vol. Biblio. of Palaeopalynology, 1836-1966. Rev. Palaeobot. Palynology.*, 8, 572 pp.
- MARCHANT, S., 1972. Evolution of the genus *Chryso-coccyx*. *Ibis*, 114, p. 219-233.
- MARGOLIS, S. V. and J. P. KENNETT, 1970. Antarctic glaciation during the Tertiary recorded in sub-Antarctic deep-sea cores. *Science*, 170(3962), p. 1085-1087.

- , 1971a. Cenozoic paleoglacial history of Antarctica recorded in subantarctic deep sea cores. *Am. Jour. Sci.*, 271(1), p. 1-36.
- , 1971b. Paleoglacial history of Antarctica recorded in deep sea cores. *Antarct. Jour.*, 6(5), p. 175-176.
- MAXWELL, A. E., R. P. VON HERZEN, K. J. HSÜ, J. E. ANDREWS, T. SAITO, S. F. PERCIVAL, Jr., E. D. MILOW, R. E. BOYCE, 1970. Deep sea drilling in the South Atlantic. *Science*, 168 (3935), p. 1047-1058.
- MATTHEW, W. D., 1915. Climate and evolution. *Ann. N. Y. Acad. Sci.*, 24, p. 171-318.
- MAYR, E., 1941. The origin and the history of the bird fauna of Polynesia. *Proc. Sixth Pacific Sci. Cong. (1939)*, 4, p. 197-216.
- , 1944a. The birds of Timor and Sumba. *Bull. Am. Mus. Nat. Hist.*, 83(art. 2), p. 127-194.
- , 1944b. Timor and the colonization of Australia by birds. *Emu*, 46, p. 113-130.
- , 1945a. *Birds of the Southwest Pacific*. Macmillan, New York, 316 pp.
- , 1945b. Wallace's Line in the light of recent zoogeographic studies. *Science and Scientists in the Netherlands Indies*, Board of the Netherlands Indies, Surinam, and Curacao, New York, p. 241-250.
- , 1963. *Animal species and evolution*. Belknap Press of Harvard Univ. Press, Cambridge, 797 pp.
- , 1972. Continental drift and the history of the Australian bird fauna. *Emu* 72(1), p. 26-28.
- MAYR, E. and D. AMADON, 1951. A classification of recent birds. *Am. Mus. Novitates*, (1946), 42 pp.
- MCDUGALL, I. and M. W. McELHINNY, 1970. The Rajmahal Traps of India—K-Ar ages and palaeomagnetism. *Earth Planet. Sci. Lett.*, 9, p. 371-378.
- McELHINNY, M. W., 1970. Formation of the Indian Ocean. *Nature*, 228, p. 977-979.
- McELHINNY, M. W. and P. WELLMAN, 1969. Polar wandering and sea-floor spreading in the southern Indian Ocean. *Earth Planet. Sci. Lett.*, 6, p. 198-204.
- MCGOWRAN, B., 1971. Unpublished report. Australia-Antarctica separation and the Eocene transgression in southern Australia. Dept. Mines. S. Australia, G.S. No. 4655.
- McINTYRE, D. J. and G. J. WILSON, 1966. Preliminary palynology of Antarctic Tertiary cratites. *New Zealand Jour. Bot.*, 4, p. 315-321.
- McKENNA, M. C., 1972a. Eocene final separation of the Eurasian and Greenland-North American landmasses. 24th Internat. Geol. Cong., Sect. 7, Paleont., Montreal, p. 275-281.
- , 1972b. Possible biological consequences of plate tectonics. *Bio. Science*, 22(9), p. 519-525.
- , 1973 (1974). Sweepstakes, filters, corridors, Noah's arks, and beached Viking funeral ships in paleogeography. In: Tarling, D. H. and S. K. Runcorn, *Implications of continental drift to the earth sciences*, Vol. 1. London, Academic Press, p. 295-308.
- McKENZIE, D. and J. G. SCLATER, 1971. The evolution of the Indian Ocean since the Late Cretaceous. *Geophys. Jour. Roy. Astr. Soc.*, 25, p. 437-528.
- MEISE, W., 1963. Verhalten der straussartigen Vögel und Monophylie der Ratitae. *Proc. Int. Ornith. Cong.*, 13, p. 115-125.
- MERCER, J. H., 1973. Cainozoic temperature trends in the southern hemisphere: Antarctic and Andean glacial evidence. In: E. M. van Zinderen Bakker, *Palaeoecology of Africa*, 8, Balkema, Cape Town, p. 85-114.
- MILLER, A. H., 1962. The history and significance of the fossil *Casuarius lydekkeri*. *Rec. Australian Mus.*, 25(10), p. 235-237.
- , 1963a. Fossil ratite birds of the late Tertiary of South Australia. *Rec. S. Australian Mus.*, 14(3), p. 413-420.
- , 1963b. The fossil flamingos of Australia. *Condor*, 65(4), p. 289-299.
- , 1966a. An evaluation of the fossil aningas of Australia. *Condor*, 68(4), p. 315-320.
- , 1966b. The fossil pelicans of Australia. *Mem. Queensland Mus.*, 14(5), p. 181-190.
- MITCHELL, T. L., 1839. Three expeditions into the interior of eastern Australia, 2. T. and W. Moore, London, 415 pp.
- NELSON, G. J., 1969. The problem of historical biogeography. *Systematic Zoo.*, 18, p. 243-246.
- NEWELL, N.D., 1971. An outline history of tropical organic Reefs. *Am. Mus. Novitates*, 2465, 37 pp.
- , 1972. The evolution of reefs. *Sci. Am.*, 226(6), p. 54-65.
- OLSON, S. L., 1974 (in press). The fossil rails of C. W. DeVis, being mainly an extinct form of *Tribonyx mortierii* from Queensland. *Emu*.
- and J. FARRAND, Jr., 1974. *Rhegminornis* restudied: a tiny Miocene turkey. *Wilson Bull.*, 86(2), p. 114-120.
- OWEN, R., 1872. Untitled. *Proc. Zool. Soc. London*, p. 682-683.
- , 1874. On *Dinornis* Pars 19: containing a description of a femur indicative of a new genus of large wingless birds (*Dromornis australis*) from a post-Tertiary deposit in Queensland. *Trans. Zool. Soc. London*, 8, p. 381-384.
- , 1879a. Memoirs on the extinct wingless birds of New Zealand, with an appendix on those in England, Australia, Newfoundland, Mauritius and Rodriguez. John van Voorst, London, vol. 1, 512 pp., vol. 2, pls. 1-130.
- , 1879b. On *Dinornis*: containing a restoration of the skeleton of *Dinornis maximus* (Owen), with an appendix on additional evidence of the genus *Dromornis* in Australia. *Trans. Zool. Soc. London*, 10(3), p. 147-188.
- PALMIERI, V., 1971. Tertiary subsurface biostratigraphy of the Capricorn Basin. *Rept. Geol. Surv. Queensland*, 52, 18 pp.
- PARKES, K. C. and G. A. CLARK, Jr., 1966. An additional character linking ratites and tinamous, and an interpretation of their monophyly. *Condor*, 68, p. 459-471.
- PASCUAL, R. and O. E. O. RIVAS, 1971. Evolucion de las comunidades de los vertebrados del Terciario Argentino, Los Aspectos Paleozoogeograficos y Paleoclimaticos relacionados. *Ameghiniana*, 7(3/4), p. 372-412.
- PETERS, J. L., 1931. Check-list of birds of the World, 1. Harvard Univ. Press, Cambridge, 345 pp.
- , 1934. Check-list of birds of the World, 2. Harvard Univ. Press, Cambridge, 401 pp.

- , 1937. Check-list of birds of the World, 3. Harvard Univ. Press, Cambridge, 311 pp.
- , 1940. Check-list of birds of the World, 4. Harvard Univ. Press, Cambridge, 291 pp.
- , 1945. Check-list of birds of the World, 5. Harvard Univ. Press, Cambridge, 306 pp.
- PITMAN, W. C., III, E. M. HERRON, and J. R. HEIRTZLER, 1968. Magnetic anomalies in the Pacific and sea-floor spreading. *Jour. Geophys. Res.*, 73(6), p. 2069-2085.
- PLANE, M. D., 1967. Stratigraphy and vertebrate fauna of the Otibanda Formation, New Guinea. *Bull. Bur. Min. Res.*, 86, 64 pp.
- PLUMSTEAD, E. P., 1962. Fossil floras of Antarctica. *Trans-Antarctic Exp. Sci. Rep.*, 9, p. 1-154.
- , 1964. Palaeobotany of Antarctica. In: Adie, R. J. (ed.), *Antarctic Geology*, North Holland Publ. Co., Amsterdam, p. 637-654.
- RAND, A. L. and E. T. GILLIARD, 1968. Handbook of New Guinea birds. Natural History Press, Garden City, New York, 628 pp.
- RAVEN, P. H. and D. I. AXELROD, 1972. Plate tectonics and Australasian Paleobiogeography. *Science*, 176(4042), p. 1379-1386.
- RENSCH, B., 1931. Die Vogelwelt von Lombok, Sumbawa und Flores. *Mitt. Zool. Mus. Berlin*, 17, p. 451-637.
- REYMENT, R. A., 1969. Ammonite biostratigraphy, continental drift and oscillatory transgressions. *Nature*, 224, p. 137-140.
- RICH, P. V., 1972. A fossil avifauna from the upper Miocene Beglia Formation of Tunisia. *Notes Serv. Geol. Tunisie*, 35, *Trav. Geol. Tunisie*, No. 5, Formation Beglia-Fasc. 1, p. 29-66.
- , 1973. The history of Australia's non-passeriform birds. Ph.D. thesis, Columbia University, New York.
- , 1974. Significance of the Tertiary avifaunas from Africa (with emphasis on a mid to late Miocene avifauna from southern Tunisia). *Annals Geol. Surv. Egypt*, 4, p. 167-210.
- RICH, P. V. and R. H. GREEN, 1974. Bird footprints at South Mt. Cameron, Tasmania. Vol. 74, part 4, of *Emu*, 1974, p. 245-248.
- RICKLERS, R. E. and G. W. COX, 1972. Taxon cycles in the West Indian avifauna. *Am. Naturalist*, 106(948), p. 195-219.
- RIDD, M. F., 1971. South-east Asia as a part of Gondwanaland. *Nature*, 234, p. 531-533.
- ROTRAMEL, G. L., 1973. The development and application of the area concept in biogeography. *Systematic Zool.*, 22(3), p. 227-232.
- SCHAEFFER, B., 1969. Mesozoic fishes and climate. *Proc. North Am. Paleont. Convention*, Pt. D, p. 376-388.
- , 1972. A Jurassic fish from Antarctica. *Am. Mus. Novitates*, 2495, 17 pp.
- SCHODDE, R. and J. H. CALABY, 1972. The biogeography of the Australo-Papuan bird and mammal faunas in relation to Torres Strait. In: Walker, D., *Bridge and barrier: the natural cultural history of Torres Strait*. *Aust. Nat. Univ. Pub. BG/3*, Canberra, p. 257-300.
- SCHOPF, J. M., 1969. Ellsworth Mountains: Position in West Antarctica due to sea floor spreading. *Science*, 164, p. 63-66.
- SCLATER, P. L., 1858. On the general geographical distribution of the members of the class Aves. *Jour. Proc. Linnean Soc. (Zool.)*, 2, p. 130-145.
- SERVENTY, D. L., 1972. Causal zoogeography Australia. *Proc. XV Internat. Ornithol. Cong.*, The Hague, p. 574-584.
- , 1973. Origin and structure of Australian bird fauna. In: Macdonald, J. D. (ed.), *Birds of Australia*, A. H. and A. W. Reed, Sydney, p. 21-28.
- SEWARD, A. C., 1941. *Plant life through the ages*. Cambridge Univ. Press, London and New York, 2nd ed., 607 pp.
- SIBLEY, C. G., 1960. The electrophoretic patterns of avian egg-white proteins as taxonomic characters. *Ibis*, 102(2), p. 215-284.
- , 1966. Protein evidence of avian relationships. *XIV Internat. Ornith. Cong., Abs.*, Oxford, p. 16-17.
- SIBLEY, C. G. and J. E. AHLQUIST, 1972. A comparative study of the egg white proteins of non-passerine birds. *Bull. Peabody Mus. Nat. Hist.*, 39, 276 pp.
- SIBLEY, C. G. and C. FRELIN, 1972. The egg white protein evidence for ratite affinities. *Ibis*, 114, p. 377-387.
- SIMPSON, E., *et al.*, 1972. Leg 25 DSDP Western Indian Ocean. *Geotimes*, 17(11), p. 21-24.
- SIMPSON, G. G., 1940. Mammals and land bridges. *Jour. Wash. Acad. Sci.*, 30(4), p. 137-163.
- , 1945. The principles of classification and a classification of mammals. *Bull. Am. Mus. Nat. Hist.*, 85, 350 + pp. xvi.
- , 1946. Fossil penguins. *Bull. Am. Mus. Nat. Hist.*, 87(art. 1), 99 pp.
- , 1953. *Evolution and geography*. Condon Lectures, Oregon State System Higher Education, Eugene.
- , 1957. Australian fossil penguins, with remarks on penguin evolution and distribution. *Rec. S. Australian Mus.*, 13, p. 51-70.
- , 1959. A new fossil penguin from Australia. *Proc. Roy. Soc. Victoria*, 71, p. 113-119.
- , 1965. New record of a fossil penguin in Australia. *Proc. Roy. Soc. Victoria*, 79, p. 91-93.
- , 1970. Miocene penguins from Victoria, Australia, and Chubut, Argentina. *Mem. Nat. Mus. Vic.*, 31, p. 17-24.
- SLATER, P. L., 1971. A field guide to Australian birds. Non-passerine. Livingstone Publ. Co., Wynnewood, Penn., 428 pp.
- SMITH, A. G. and A. HALLAM, 1970. The fit of Southern continents. *Nature*, 225, p. 139-144.
- SOWERBUTTS, W. T. C., 1972. Rifting in eastern Africa and the fragmentation of Gondwanaland. *Nature*, 235, p. 435-437.
- SPENCER, B. and J. A. KERSHAW, 1910. A collection of sub-fossil bird and marsupial remains from King Island, Bass Strait. *Mem. Nat. Mus. Victoria*, 3, p. 5-35.
- STAUFFER, P. H. and D. J. GOBBETT, 1972. Southeast Asia a part of Gondwanaland? *Nature, Phys. Sci.*, 240, p. 139.
- STEVENS, G. R. and R. N. CLAYTON, 1971. Oxygen isotope studies on Jurassic and Cretaceous belemnites from New Zealand and their biogeographic significance. *N.Z. Jour. Geol. Geophys.*, 14(4), p. 829-897.

- STIRLING, E. C., 1896. The newly discovered extinct gigantic bird of South Australia. *Ibis*, 7(11), p. 593.
- STIRLING, E. C. and A. H. C. ZIETZ, 1896. Preliminary notes on *Genyornis newtoni*; a new genus and species of fossil struthious bird found at Lake Callabonna, South Australia. *Trans. Proc., Roy. Soc. Australia*, 20, p. 171-190.
- , 1900. Fossil remains of Lake Callabonna: 1. *Genyornis newtoni*. A new genus and species of fossil struthious bird. *Mem. Roy. Soc. S. Australia*, 1(2), p. 41-80.
- , 1905. Fossil remains of Lake Callabonna, Pt. III. Description of the vertebrae of *Genyornis newtoni*. *Mem. Roy. Soc. S. Australia*, 1(pt. 3), p. 81-110.
- , 1913. Fossil remains of Lake Callabonna. Pt. IV. Description of some further remains of *Genyornis newtoni*. *Mem. Roy. Soc. S. Australia*, 1(4), p. 111-126.
- STIRTON, R. A., R. H. TEDFORD, and M. O. WOODBURN, 1967. A new Tertiary formation and fauna from the Tirari Desert, South Australia. *Rec. S. Australia, Mus.*, 15(3), p. 427-462.
- , 1968. Australian Tertiary deposits containing terrestrial mammals. *Univ. Calif. Publ. Geol. Sci.*, 77, 30 pp.
- STORER, R. W., 1960. The classification of birds. In: Marshall, A. J. (ed.), *Biology and comparative physiology of birds*, 1, Academic Press, London, p. 57-93.
- , 1971. Classification of birds. In: Farner, D. S. and J. R. King (ed.), *Avian biology*, 1, Academic Press, New York, p. 1-18.
- STRESEMANN, E., 1927-1934. Geographische Verbreitung. In: Kükenenthal, W., *Handb. Zool.*, 7(2), Aves, p. 633-658.
- , 1939-41. Die Vögel von Celebes. *Jour. Ornith.*, 87, p. 312-425; 88, p. 1-135, 389-487; 89, p. 1-102.
- SZALAY, F. S. and M. C. MCKENNA, 1971. Beginning of the age of mammals in Asia: the late Paleocene Gashato fauna, Mongolia. *Bull. Am. Mus. Nat. Hist.*, 144(4), p. 271-317.
- TALENT, J. A., P. M. DUNCAN, and P. L. HANDLEY, 1966. Early Cretaceous feathers from Victoria. *Emu*, 66(2), p. 81-86.
- TARLING, D. H., 1971. Gondwanaland, palaeomagnetism and continental drift. *Nature*, 229, p. 17-21, 71.
- TARLING, D. and M. TARLING, 1971. *Continental drift. A study of the earth's moving surface.* Doubleday and Co., Inc., Garden City, New York, 140 pp.
- TEDFORD, R. H., 1973a (in press). Marsupials and the new biogeography. S.E.P.M. Symposium, Paleogeographic provinces and provinciality, Denver, 1972, ed. C. A. Ross.
- , 1973b (ms.). Preliminary Report on collections of the American Museum of Natural History, 1971. Report to the South Australian Dept. of Mines, 1973.
- VALENCIO, D. A. and J. F. VILAS, 1972. Palaeomagnetism of late Paleozoic and early Mesozoic rocks of South America. *Earth Planet. Sci. Lett.*, 15(1), p. 75-84.
- VAN BEMMELEN, R. W., 1949. *The geology of Indonesia*, 1. Gov't. Print. Office, The Hague, 732 pp.
- VAN DER LINDEN, W. J. M., 1969. Rotation of the Melanesian complex and of West Antarctica—a key to the configuration of Gondwana? *Palaeogeog., Palaeoclim., Palaeoecol.*, 6, p. 37-44.
- VAN TETS, G. F., 1974a. A revision of the fossil Megapodiidae, including a description of a new species of *Progura* DeVis. *Trans. Roy. Soc. S. Australia*, 98(4), p. 213-224.
- , 1974b. Fossil birds (Aves) from Weeke's Cave, Nullarbor Plain, South Australia, 98(4), p. 229-230.
- , 1974c. Was '*Taphaetus*' *lacertosus* DeVis a fishing eagle, *Ichthyophaga* Lesson? *Emu*, 74, p. 58.
- VAN TETS, G. F. and M. J. SMITH, 1974. Small fossil vertebrates from Victoria Cave, Naracoorte, South Australia III. Birds (Aves). *Trans. Roy. Soc. S. Australia*, 98(4), p. 225-228.
- VEEVERS, J. J., 1971. Phanerozoic history of Western Australia related to continental drift. *Jour. Geol. Soc. Australia*, 18(2), p. 87-96.
- and Evans, 1973. Sedimentary and magnetic events in Australia and the mechanism of worldwide Cretaceous transgressions. *Nature Phys. Sci.*, 245(142), p. 33-36.
- VEEVERS, J. J., J. G. JONES, and J. A. TALENT, 1971. Indo-Australian stratigraphy and the configuration and dispersal of Gondwanaland. *Nature*, 229, p. 383-388.
- VERHEYEN, R., 1953. Bijdrage tot de osteologie en die systematiek der Anseriformes. *Le Gerfaut*, 43, Suppl., p. 373-497.
- , 1955. La systématique des Anseriformes basée sur l'ostéologie comparée. *Bull. Inst. Roy. Sci. Natur. Belgique*, 31, p. 35-38.
- VILAS, J. F. and D. A. VALENCIO, 1970. The recurrent Mesozoic drift of South America and Africa. *Earth Planet. Sci. Lett.*, 7, p. 441-444.
- VINE, F. J. and D. H. MATTHEWS, 1963. Magnetic anomalies over ocean ridges. *Nature*, 199(4897), p. 947-949.
- VOGT, P. R. and J. R. CONOLLY, 1971. Tasmanian guyots, the age of the Tasman Basin, and motion between the Australian plate and the mantle. *Bull. Geol. Soc. Amer.*, 82(9), p. 2577-2584.
- VON DER BORCH, C., et al., 1972. Deep sea drilling project, Leg 22. *Geotimes*, 17(6), p. 15-17.
- WALDMAN, M., 1970. A third specimen of a lower Cretaceous feather from Victoria, Australia. *Condor*, 72, p. 377.
- WALKER, E. P., et al., 1964. *Mammals of the World*, 2. Johns Hopkins Press, Baltimore, p. 647-1500.
- WALLACE, A. R., 1863. A list of the birds inhabiting the islands of Timor, Flores, and Lombok, etc. *Proc. Zool. Soc. London*, p. 480-497.
- , 1869. *The Malay Archipelago.* Harper and Bros., New York, 638 pp.
- , 1876. *Geographical distribution of animals*, 1. Macmillan and Co., London, 503 pp.
- WEISSEL, J. K. and D. E. HAYES, 1972. Magnetic anomalies in the southeast Indian Ocean. In Hayes, D. E. (ed.), *Antarctic Oceanology II. The Australian-New Zealand sector.* Amer. Geophy. Union, Antarctic Res. Ser. Mon., 19, p. 165-196.
- WELLMAN, P. and M. W. McELHINNY, 1970. K-Ar age of the Deccan Traps, India. *Nature*, 227, p. 595-596.

- WELLMAN, P., M. W. MCELHINNY, and I. MCDUGALL, 1969. On the polar-wander path for Australia during the Cenozoic. *Geophy. Jour. Roy. Astr. Soc.*, 18, p. 371-395.
- WILKINSON, H. E., 1969. Description of an upper Miocene albatross from Beaumaris, Victoria, Australia, and a review of fossil Diomedidae. *Mem. Natn. Mus. Victoria*, 29, p. 41-51.
- WOOLFENDEN, G. E., 1961. Postcranial osteology of the waterfowl. *Florida State Mus., Biol. Sci.*, 6(1), p. 1-129.